

## **REMARKS/ARGUMENTS**

### **Status of Claims**

Claims 31, 32, 34, and 36-37 are pending and are under examination. Claims 1-30, 33, and 35 are cancelled.

### **Amendments to the Claims**

No amendments to the claims were made.

### **Priority of Claims 31, 32, 34, and 36-37**

This application claims benefit of U.S. provisional application no. 60/491,350 ("350 Application"), filed July 31, 2003 and claims benefit of U.S. provisional application no. 60/509,037 ("037 Application") filed October 4, 2002 (converted from non-provisional application no. 10/264,825).

### **Sequence Compliance**

Applicants acknowledge the Examiner's withdrawal of the objection to the sequence listing.

### **Specification**

Applicants acknowledge the Examiner's withdrawal of the objection to the specification.

### **Withdrawal of Rejections**

Applicants acknowledge the Examiner's withdrawal of the rejections made under 35 U.S.C. § 112, second paragraph and 35 U.S.C. § 112, first paragraph. As such, the instant application provides written description and enablement for pending claims 31, 32, 34, and 36-37.

**Drawings**

The Examiner did not acknowledge the drawings which were originally filed. Specifically, the Examiner did not indicate in the Office Actions mailed August 22, 2006 and May 1, 2007 whether the drawings submitted by Applicants were accepted or objected to by the Examiner. Applicants respectfully request acknowledgement of the acceptance of the drawings or objection by checking the appropriate box in the next Office Action.

**Applicants' Invention**

Applicants discovered that cancer cells overexpress a protein, Dvl-3, and that inhibiting expression of Dvl-3 inhibits the growth of cancer cells overexpressing Dvl-3. Nothing in the prior art suggested this invention.

**Claim Rejection - 35 USC § 102(b)**

The Examiner rejected Claims 31 and 37 under 35 U.S.C. § 102(b) as being anticipated by Song *et al.* (*J. Biol. Chem.* 275:23790-23797 (2000); "Song"). According to the Examiner, Song teaches that (i) protein kinase CK2 is involved in tumorigenesis (p. 23790, col. 2), (ii) CK2 is important to modulate phosphorylation of Dvl-3 which is expressed in breast cancer cells because when breast cancer cells were treated with apigenin, a CK2 inhibitor, the phosphorylation of Dvl-3 protein is diminished (Figure 5), (iii) apigenin reduces the levels of Dvl-3 protein in breast cells, and (iv) apigenin inhibits cell proliferation.

The rejection is respectfully traversed.

**A. The Legal Standard**

For a rejection of claims under §102 to be properly founded, the Examiner must establish that a single prior art reference either expressly or inherently discloses each and every element of the claimed invention. *See, e.g. Hybritech Inc. v. Monoclonal Antibodies, Inc.*, 231 USPQ 81 (Fed. Cir. 1986), *cert denied*, 480 U.S. 947 (1987); and *Verdegaal Bros. V. Union Oil Co. of California*, 2 USPQ2d 1051, 1053 (Fed. Cir. 1987). In *Scripps Clinic & Research Found. V. Genentech, Inc.*, 18 USPQ2d 1001 (Fed. Cir. 1991), the Federal Circuit held that:

"Invalidity for anticipation requires that **all of the elements and limitations** of the claim are found **within a single prior art reference**.... There must be no difference between the claimed invention and the reference disclosure, as viewed by a person of ordinary skill in the field of the invention." *Id.* at 1010.

Anticipation cannot be found, therefore, unless a cited reference discloses all of the elements, features or limitations of the presently claimed invention. Applicants respectfully submit that Song fails to recite all of the elements of claims 31 and 37.

**B. Song Does Not Disclose A Cancer Cell That Overexpresses A Dvl-3 Protein And Does Not Disclose An Agent That Inhibits Dvl-3 Expression**

The Examiner acknowledged that nowhere does Song teach a cancer cell (claim 31) or a breast cancer cell (claim 37) *overexpressing* a Dvl-3 protein. Song does not compare a normal cell and a cancer cell or a normal cell and a breast cancer cell to determine that the cancer cell or breast cancer cell overexpresses a Dvl-3 protein. Further, Applicants submit that, contrary to the Examiner's allegation, Figure 5 of Song does not show that apigenin, an inhibitor of CK2, diminishes phosphorylation of Dvl-3. Only Figure 5B reports an experiment using apigenin and this particular experiment refers to the diminished phosphorylation of  $\beta$ -catenin, not Dvl-3 (Figure 5B).

Song does not teach or suggest an agent that inhibits Dvl-3 *expression*. Song teaches that apigenin, an inhibitor of CK2, through an unknown mechanism, causes the degradation of a Dvl-3 protein that *already exists* in a cell. Song does not teach or suggest that apigenin inhibits Dvl-3 *expression* (i.e., transcription of a Dvl-3 mRNA from a Dvl-3 encoding gene or translation of the Dvl-3 mRNA to produce a Dvl-3 protein, as one of ordinary skill in the art would understand the term "expression" in the context of Applicants' claims). Song states on page 23795, col. 1 in the context of Figures 6 and 7:

"To determine whether the reduction in  $\beta$ -catenin occurred through a decreased rate of synthesis or increased rate of degradation, we measured the half-life of the protein in the presence of a *cycloheximide* that blocked new protein synthesis. We found that  $\beta$ -catenin is quite stable in Wnt-1-expressing cells, with a half-life of more than 5 h (Fig. 7) ... The *Dvl*

*proteins appear to be equally stable. However, in the presence of apigenin, protein levels rapidly decline. Immunoreactive Dvl proteins disappears in less than 30 min..."* (emphasis added)

Because cycloheximide is a protein synthesis inhibitor that acts specifically on the 60S subunit of eukaryotic ribosomes, Song investigated the effect of apigenin *on already expressed* Dvl-3 protein and did not investigate the effect of an agent that inhibited the expression of Dvl-3 protein.

Claims 31 and 37 require an agent that inhibit the *expression* of a Dvl-3 protein. Song does not teach or suggest all limitations of claims 31 and 37. Therefore, Song does not anticipate claims 31 and 37.

Applicants respectfully request withdrawal of the rejection of these claims under 35 U.S.C. § 102(b).

#### **Claim Rejection - 35 USC § 102(e)**

The Examiner rejected Claims 31, 32, 34, and 37 under 35 U.S.C. § 102(e) as being anticipated by Alsobrook *et al.* (US 20030229016 based on U.S. Application Ser. No. 10/307,928 ("928 Application"), filed December 2, 2002 and published December 11, 2003; priority to 8/26/02 and earlier; "Alsobrook"). According to the Examiner, Alsobrook teaches methods for treating a cancer cell such as a lung cancer cell or breast cancer cell [0016] using an siRNA [0080] which inhibits expression of a splice variant of a dishevelled-3-like protein (Table 1). The Examiner appreciated that Applicants' claims are not limited to any kind of Dvl-3 protein moiety and argued that Alsobrook's disclosure of the use of the splice variant for a Dvl-3 like protein for inhibiting the expression of the Dvl-3 like protein to inhibit cancer cell proliferation (i.e., using siRNA as an agent) anticipates Applicants' claims 31, 32, 34, and 37.

The rejection is respectfully traversed.

#### **A. The Legal Standard**

The legal standard for a rejection of claims under §102 is discussed *supra*.

**B. None Of The Alsobrook Earlier Filed Provisional Patent Applications Disclose The Subject Matter Of Applicants' Claims**

Alsobrook was filed as U.S. Application Ser. No. 10/307,928 ("928 Application") on December 2, 2002, claiming priority to eleven (11) provisional patent applications, including:

- (1) 60/406,353 ("353 Application"), filed August 26, 2002;
- (2) 60/401,788 ("788 Application"), filed August 7, 2002;
- (3) 60/384,024 ("024 Application"), filed May 29, 2002;
- (4) 60/383,744 ("744 Application"), filed May 28, 2002;
- (5) 60/381,495 ("495 Application"), filed May 17, 2002;
- (6) 60/380,981 ("981 Application"), filed May 15, 2002;
- (7) 60/373,288 ("288 Application"), filed April 17, 2002;
- (8) 60/344,903 ("903 Application"), filed December 31, 2001;
- (9) 60/342,592 ("592 Application"), filed December 20, 2001;
- (10) 60/341,540 ("540 Application"), filed December 17, 2001; and
- (11) 60/341,477 ("477 Application"), filed December 17, 2001, collectively

referred to as "Alsobrook provisional applications."

To the extent that these Alsobrook provisional applications were available on PAIR for Applicants' review, Applicants submit that none of these Alsobrook provisional applications teaches a method for treating a cancer cell (such as a lung cancer or breast cancer cell) that *overexpresses* a Dvl-3 protein by contacting the cell with an agent (such as a siRNA) that inhibits Dvl-3 expression wherein the growth of the cancer cell is inhibited.

None of the '353, '788, '024, '744, '495, '981, '288, '592, '540, and '477 Applications disclose the Dvl-3 splice variant. The '903 Application, filed on December 31, 2001, discloses on pages 53 to 79 a Dvl-3 splice variant which is also disclosed in Alsobrook's '928 Application. The remainder of the '903 Application, however, includes disclosure which is unrelated to Dvl-3, but rather discloses proteins and nucleic acids for Colonic And Hepatic Tumor Over-Expressed Protein-like Proteins, Acetyltransferase-like Proteins, Granzyme H-like Proteins, Fibulin-2-like Proteins, 4930418P06RIK Rhomboid-like Proteins, DORA Protein Precursor-like Proteins, IPAS-like Proteins, splice variants of Cartilage Oligomeric Matrix

Protein-like Proteins, and splice variants of Insulin-like Growth Factor Binding Protein 4 (IGFBP4)-like Proteins.

Pages 53 to 79 of the '903 Application are provided for the Examiner's review as **Exhibit A**. Applicants submit that with respect to the disclosure of the Dvl-3 splice variant, the '903 Application discloses various sequence alignments, hydropathy data (Figures 1-5), and tissue expression data of Dvl-3 (page 59). Specifically, with respect to expression of the Dvl-3 splice variant, the '903 Application discloses expression of the Dvl-3 splice form in various normal tissues and two tumors (ovary and parathyroid gland) (page 59).

Applicants submit that the '903 Application, however, does not disclose all limitations of Applicants' claims 31, 32, 34, and 37. For example, the '903 Application does not teach *overexpression* of the Dvl-3-like protein in any cancer cell, such as a lung cancer cell or breast cancer cell. In fact, the '903 Application does not even mention lung cancer or breast cancer. The '903 Application does also not disclose an agent, such as an siRNA, for the inhibition of Dvl-3 expression.

As such, Alsobrook is not entitled to benefit of the priority date of the '903 Application for allegedly disclosing Applicants' subject matter of claims 31, 32, 34, and 37.

Because none of the other Alsobrook provisional applications provides the a disclosure of the Dvl-3 splice variant, Alsobrook is also not entitled to claim benefit of any of these Alsobrook provisional patent applications for the alleged disclosure of the subject matter of Applicants' claims 31, 32, 34, and 37.

As such, in rejecting claims 31, 32, 34, and 37 under 102(e) as allegedly being anticipated by Alsobrook, the Examiner must rely on the disclosure of the '928 Application, which has a filing date of December 2, 2002.

**C. Alsobrook's '928 Application Does Not Qualify As Prior Art Under 35 U.S.C. § 102(e)**

The presently examined application claims benefit of U.S. Provisional Application No. 60/509,037 ("037 Application") filed October 4, 2002. This filing date predates the filing date of Alsobrook's '928 Application by two months. Specifically, Applicants' claim

31 is supported by the '037 Application (see, for example, page 17, lines 20-21; page 36, lines 23-24; page 37, lines 29-32, page 38, lines 10-15, page 38, lines 20-21; Figure 9). Therefore, the '928 Application does not qualify as prior art under 35 U.S.C. § 102(e) and the rejection of claim 31 should be withdrawn. Claims 32, 34, and 37 depend on claim 31 and incorporate the limitations of claim 31. Thus, Alsobrook is also not prior art against dependent claims 32, 34, and 37.

Applicants respectfully request withdrawal of the rejection of claims 31, 32, 34, and 37 under 35 U.S.C. § 102(e).

**D. Alsobrook's '928 Application Does Not Disclose A Cancer Cell That Overexpresses A Dvl-3 Protein**

As discussed above, the only potentially relevant Alsobrook application is the '928 Application. Applicants submit that the '928 Application does not anticipate Applicants' claims because it does not teach all of the limitations of the claims.

Alsobrook does not teach or suggest a cancer cell (claims 31, 34), a lung cancer cell (claim 32), or a breast cancer cell (claim 37) *overexpressing* a Dvl-3 protein. Thus, Alsobrook does not teach the limitation of Applicants' claim "a cancer cell that overexpresses a Dvl-3 protein." As such Alsobrook does not teach all limitations of Applicants' claims and it is an improper §102 reference.

Applicants respectfully request withdrawal of the rejection of claims 31, 32, 34, and 37 under 35 U.S.C. § 102(e).

**E. Applicants' Invention Predates Alsobrook's '928 Application**

The Alsobrook '928 Application is cited by the Examiner as a 102(e) reference. It is, thus, subject to swearing behind. Accordingly, without conceding that Alsobrook's '928 Application provides an enabling disclosure of each and every element and limitation for the subject matter of Applicants' claims 31, 32, 34, and 37, Applicants herewith submit a Declaration under 37 CFR 1.131 which establishes that Applicants completed their invention prior to the effective filing date of Alsobrook's '928 Application, which is December 2, 2002. Evidence of

Applicants' conception of the invention prior to December 2, 2002 includes (i) the finding that tumor cells when compared to normal cells overexpress Dvl-3 mRNA; (ii) the finding that cancer cells, including lung cancer cells, breast cancer cells and mesothelioma, overexpress a Dvl-3 protein when compared to normal or non-tumor cells; (iii) designing Dvl-3 siRNA nucleic acids for inhibition of Dvl-3 expression; and (iv) ordering Dvl-3 siRNA nucleic acids for inhibition of Dvl-3 expression. After conceiving of the invention, Applicants diligently worked towards actual and constructive reduction to practice their invention.

In view of the arguments provided herein and further in view of the Rule 131 Declaration, Alsobrook is no longer considered anticipatory art. Applicants submit that the rejection of claims 31, 32, 34, and 47 over Alsobrook has been fully addressed. Reconsideration and withdrawal of this reference as a basis for the 35 U.S.C. §102(e) rejection is respectfully requested.

**Claim Rejection - 35 USC § 103(a)**

**A. The Legal Standard**

Establishing a *prima facie* case for obviousness under § 103 requires the Examiner show, *inter alia*:

- (1) The prior art references teach or suggest all claim limitations of the rejected claim(s). *In re Royka*, 180 USPQ 580 (CCPA 1974); and MPEP §2143.03.
- (2) The existence of some suggestion or motivation, either in the references themselves or in the knowledge generally available to one of ordinary skill in the art, to modify the reference or to combine the reference teachings. *In re Fine*, 837 F.2d 1071, 5 USPQ2d 1596 (Fed. Cir., 1988).
- (3) A reasonable expectation of success in combining the references. This must be found in the prior art, and not in the applicants' disclosure. *In re Vaeck*, 947 F.2d 488, 20 USPQ2d 1438 (Fed. Cir., 1991).

A *prima facie* case of obviousness requires the Examiner to provide an explicit reason why one of ordinary skill in the art would combine the known elements in the fashion claimed by Applicants. Recently, in reviewing this standard, the Supreme Court noted that any



analysis supporting a rejection under § 103(a) must be made explicit, and that it is "important to identify a reason that would have prompted a person of ordinary skill in the relevant field to combine the [prior art] elements in the manner claimed. *KSR Intl Co. v. Teleflex Inc.*, 82 USPQ2d 1385, 1396 (U.S. 2007). "This is so because inventions in most, if not all, instances rely upon building blocks long since uncovered, and claimed discoveries almost of necessity will be combinations of what, in some sense, is already known." *Id.* To support a rejection under § 103 using the Federal Circuit's teaching-suggestion-motivation (TSM) test, the Office must provide evidence that demonstrates some suggestion or motivation to modify or combine the references, whether in the references themselves or in the knowledge generally available to one of ordinary skill in the art. *In re Fine* 837 F.2d at 1074, MPEP § 2143.

A *prima facie* case of obviousness requires the Examiner to show that one of ordinary skill in the art would have had a reasonable expectation of success in modifying the prior art references, or in combining their relevant teachings. *In re Vaeck*, 947 F.2d 488, 493 (Fed. Cir., 1991). The teaching or suggestion to make the claimed combination *and* the reasonable expectation of success must both be found in the prior art, and *not* based on applicant's disclosure. *Id.* The Examiner's suggestion of the desirability of doing what the inventor has done must be found either expressly or impliedly in the references, or supported by a convincing line of reasoning, which must rely on logic and sound scientific reasoning. *Ex parte Clapp*, 227 USPQ 972, 973 (Bd. Pat. App. & Inter. 1985). *See also* MPEP § 2144; and *Ex parte Levengood*, 28 USPQ2d 1300 (Bd. Pat. App. & Inter. 1993) (requiring reliance on logic and sound scientific reasoning in supporting a conclusion of obviousness).

**B. Rejection of Claims 31 and 37 Over Song and Bui**

The Examiner rejected Claims 31 and 37 under 35 U.S.C. § 103(a) as being unpatentable over Song further in view of Bui *et al.* (*Biochem. Biophys. Res. Comm.* 239:510-516 (1997); "Bui"). According to the Examiner, a method of inhibiting the growth of a cancer cell, such as a breast cancer cell, with an agent that effects Dvl-3 expression was *prima facie* obvious at the time of the invention over Song and Bui. The Examiner acknowledged that Song does not teach Dvl-3 expression in cancer cells and cites to Bui to provide this teaching.

The rejection is respectfully traversed.

1. ***The Combination Of Song And Bui Fails To Teach All Elements Of the Applicants' Invention***

The teaching of Song has been discussed in detail *supra*. As also acknowledged by the Examiner (page 8 of the Office Action), Song does not disclose a cancer cell that expresses or overexpresses a Dvl-3 Protein. Further, as discussed *supra*, Song does not disclose an agent that inhibits Dvl-3 expression.

Bui merely discloses Dvl-3 expression in various cancer cell lines, including breast cancer cells, but does not disclose "an agent that inhibits Dvl-3 expression." As such, neither Song nor Bui disclose "an agent that inhibits Dvl-3 expression to inhibit the growth of a cancer cell." Bui cannot provide the missing claim element and claim limitation that is also missing in Song. Therefore, the combination of Song and Bui does not disclose all elements and all claim limitations of Applicants' claims 31 and 37.

As the combination of references suggested in the Office Action *fails to provide all of the elements* of Applicants' claimed invention, a *prima facie* case of obviousness has not been set forth. Therefore, Applicants respectfully request the rejection of claims 31 and 37 be withdrawn.

2. ***Bui Teaches Away From Applicants' Invention, There Is No Motivation To Make The Suggested Combination And No Reasonable Expectation Of Success***

Because Bui teaches that Dvl-3 is not overexpressed in cancer cells, it provides a reason against combination with Song. Therefore, one of ordinary skill in the art would not be motivated to combine the Song and Bui references. Specifically, Bui teaches in the abstract that:

"Statistically, there was no difference in DVL-3 mRNA level between normal breast tissues and tumors. In human colorectal samples, DVL-3 was expressed equally in matched normal tissues, polyps and tumors." (Emphasis added).

and on page 515, column 1:

"We have also investigated a potential role for DVL-3 in human breast and colon tumorigenesis ... Since the Wnt gene is an upstream signal of

DVL in the wingless signaling pathway, it was thought that aberrant expression of Wnt could alter DVL expression. However, the data presented here showed no difference in DVL-3 mRNA expression between normal breast tissues and corresponding tumours, and between matched normal colon tissues, polyps and tumors." (Emphasis added).

This is directly opposed to Applicants' discovery and claimed invention. Contrary to Bui's teaching, Applicants' invention requires a cancer cell to *overexpress* a Dvl-3 protein. Bui expressly teaches away from Applicants' invention. Teaching away is a strong motivation for one of ordinary skill in the art to not combine references and has been acknowledged to be strong evidence for the invention in question to be not obvious. Because of Bui's teaching away, there can also be no reasonable expectation of success in combining the Song and Bui references. The reasonable expectation of success must be found in the prior art, and not in the Applicants' disclosure.

As the references suggested in the Office Action *fail to motivate to make the suggested combination* of Applicants' claimed invention and *fail to provide a reasonable expectation of success*, Applicants' invention cannot be obvious in view of the cited art. A *prima facie* case of obviousness has not been set forth. Therefore, Applicants respectfully request the rejection of Claims 31 and 37 under 35 U.S.C. § 103(a) be withdrawn.

**C. Rejection of Claims 31 and 32 Over Song and Engelmann**

The Examiner rejected Claims 31 and 32 under 35 U.S.C. § 103(a) as being unpatentable over Song further in view of Engelmann *et al.* (*Phytomedicine* 9(6):489-495 (2202); "Engelmann"). According to the Examiner, a method of inhibiting a lung cancer cell with an agent that effects Dvl-3 expression was *prima facie* obvious at the time of the invention over Song and Engelmann. The Examiner acknowledged that Song does not teach using the method of inhibiting cancer cell growth in a lung cancer cell and cites to Engelmann to provide this teaching.

The rejection is respectfully traversed.

1. ***The Combination Of Song And Engelmann Fails To Teach All Elements Of the Applicants' Invention***

The teaching of Song has been discussed in detail *supra*. As acknowledged by the Examiner (page 8 of the Office Action), Song does not disclose a cancer cell that expresses or overexpresses a Dvl-3 Protein (claim 31) or using the method of inhibiting growth of a lung cancer cell (claim 32) (page 9 of the Office Action). Further, as discussed *supra*, Song does not disclose an agent that inhibits Dvl-3 expression.

According to the Examiner, Engelmann discloses in the abstract inhibition of lung cancer, glioma and colon cancer in vivo with apigenin.

As discussed *supra* and as acknowledged by the Examiner, apigenin is an inhibitor of CK2. While apigenin may or may not have a direct or indirect effect on Dvl-3 protein levels or protein stability as alleged by the Examiner, both Song and Engelmann references fail to provide evidence that apigenin is an agent that inhibits Dvl-3 *expression*. As such, both references taken individually or combined fail to provide an agent that inhibits Dvl-3 expression as required by Applicants' Claims 31 and 32.

As the combination of references suggested in the Office Action *fails to provide all of the elements* of Applicants' claimed invention, a *prima facie* case of obviousness has not been set forth. Therefore, Applicants respectfully request the rejection of Claims 31 and 32 be withdrawn.

2. ***There Is No Motivation To Make The Suggested Combination And No Reasonable Expectation Of Success***

There is nothing in Engelmann that would lead one of ordinary skill in the art make believe that the teaching of Engelmann would be useful for inhibiting Dvl-3 expression. Engelmann does not even mention Dvl-3. Therefore, one of ordinary skill in the art would not be motivated to combine the teaching of Song and Engelmann.

As the references suggested in the Office Action *fail to motivate to make the suggested combination* of Applicants' claimed invention and *fail to provide a reasonable expectation of success*, Applicants' invention cannot be obvious in view of the cited art. A *prima*

*facie* case of obviousness has not been set forth. Therefore, Applicants respectfully request the rejection of Claims 31 and 32 under 35 U.S.C § 103(a) be withdrawn.

**D. Rejection Of Claims 31 and 36 Over Song And You As Evidenced By Uematsu**

The Examiner rejected Claims 31 and 36 under 35 U.S.C. § 103(a) as being unpatentable over Song further in view of You *et al.* (*Proc. Am. Assoc. Cancer Res.* 42:609 (2001); "You") as evidenced by Uematsu *et al.* (*Oncogene* 22:7218-7221 (2003); "Uematsu"). According to the Examiner, a method of inhibiting a mesothelioma cell with an agent that effects Dvl-3 expression was *prima facie* obvious at the time of the invention over Song and You as evidenced by Uematsu. The Examiner acknowledged that Song does not teach using the method of inhibiting cancer cell growth in a mesothelioma and cites to You to provide this teaching.

The rejection is respectfully traversed.

**1. The Combination Of Song And You Fails To Teach All Elements Of the Applicants' Invention**

The teaching of Song has been discussed in detail *supra*. As acknowledged by the Examiner (page 8 of the Office Action), Song does not disclose a cancer cell that expresses or overexpresses a Dvl-3 Protein (claim 31) or using the method of inhibiting growth of a mesothelioma (claim 36) (page 10 of the Office Action). Further, as discussed *supra*, Song does not disclose an agent that inhibits Dvl-3 expression.

According to the Examiner, You discloses in the abstract overexpression of Dvl and its apparent involvement in inducing tumorigenicity by a canonical Wnt signaling pathway.

As the Examiner is aware, Dvl proteins include Dvl-1, Dvl-2, and Dvl-3 proteins. Applicants submit that while You discloses that a Dvl protein is overexpressed in mesothelioma cells, the abstract by You does not disclose that the Dvl protein is Dvl-3 as required by Applicants' claims. Later experiments, e.g., those disclosed in Applicants' '037 Application showed that the Dvl protein overexpressed in mesothelioma cells, as described by You, includes a Dvl-3 protein.

Further, and more importantly, You does not teach an agent that inhibits Dvl-3 expression (or Dvl expression) leading to inhibition of the growth of a cancer cell. Thus, both Song and You references taken individually or combined fail to provide an agent that inhibits Dvl-3 expression as required by Applicants' Claims 31 and 36. Therefore, both Song and You fail to provide all elements and limitations of Applicants' claims.

As the combination of references suggested in the Office Action *fails to provide all of the elements* of Applicants' claimed invention, a *prima facie* case of obviousness has not been set forth. Therefore, Applicants respectfully request the rejection of Claims 31 and 36 under 35 U.S.C § 103(a) be withdrawn.

2. ***There Is No Motivation To Make The Suggested Combination And No Reasonable Expectation Of Success***

As discussed *supra*, there is nothing in the Song and You references that teach a method for inhibiting the growth of a cancer cell with an agent for inhibiting Dvl-3 expression and achieving inhibition of the growth of the cancer cell. Therefore, one of ordinary skill in the art would not be motivated to combine the teaching of Song and You to arrive at Applicants' invention.

As the references suggested in the Office Action *fail to motivate to make the suggested combination* of Applicants' claimed invention and *fail to provide a reasonable expectation of success*, Applicants' invention cannot be obvious in view of the cited art. A *prima facie* case of obviousness has not been set forth. Therefore, Applicants respectfully request the rejection of Claims 31 and 36 under 35 U.S.C § 103(a) be withdrawn.

3. ***Without The Benefit Of Impermissible Hindsight, The Claimed Invention Was Not Obvious At The Time It Was Invented***

In *KSR*, the Court also cautioned against the use of impermissible hindsight. *KSR* at 1742. ("A factfinder should be aware, of course, of the distortion caused by hindsight bias and must be cautious of arguments reliant upon *ex post* reasoning."). Applicants respectfully submit that, without the teachings of the instant specification, one of skill in the art would not have known at the time the invention was made to inhibit the growth of a cancer cell overexpressing a

Dvl-3 protein with an agent that inhibits Dvl-3 expression. The presently claimed method provides a new way of inhibiting the growth of a cancer cell that is not suggested by the Song and/or You references. Identifying the claimed invention in a publication ("Uematsu") which was published in the journal *Oncogene* on October 16, 2003 by the inventive group of the instant application (Applicants He, You, Xu, and Jablons) after the filing date of the instant application and after its effective filing date, to allegedly fit the elements of the claims requires hindsight provided by the claimed invention. This, as emphasized in both the case law and the MPEP, is impermissible. Further, as declared in the accompanying 131 Declaration, Kazutsugu Uematsu, the first author of "Uematsu" was a post-doctoral fellow in Applicants' laboratory who worked under the supervision of Applicants.

Therefore, Applicants respectfully request the rejection of Claims 31 and 36 under 35 U.S.C § 103(a) be withdrawn.

**E. Rejection of Claims 31 and 36 Over Alsobrook And You As Evidenced by Uematsu**

The Examiner rejected Claims 31 and 36 under 35 U.S.C. § 103(a) as being unpatentable over Alsobrook in view of You as evidenced by Uematsu. According to the Examiner, a method of inhibiting a mesothelioma cell with an agent that effects Dvl-3 expression was *prima facie* obvious at the time of the invention over Alsobrook in view of You. The Examiner acknowledged that Alsobrook does not teach using the method of inhibiting cancer cell growth in a mesothelioma and cites to You to provide this teaching.

The rejection is respectfully traversed.

As an initial matter, in view of the rule 131 Declaration, Alsobrook does not qualify as prior art under 35 U.S.C. § 102(e). On this basis alone, this rejection under 35 U.S.C. 103(a) should be withdrawn.

The shortcomings of the teachings of You, Alsobrook's '928 Application, and Alsobrook's provisional patent applications have been discussed *supra*. In view of the arguments provided herein *supra* and because Alsobrook does not qualify as prior art under § 102(e), this

rejection should be withdrawn. Reciting to Uematsu, as discussed, *supra*, constitutes impermissible hindsight.

As such, the Examiner did not present a *prima facie* case of obviousness. Applicants respectfully request the rejection of Claims 31 and 36 under 35 U.S.C § 103(a) as being unpatentable over Alsobrook in view of You as evidenced by Uematsu be withdrawn.

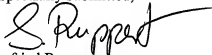
### CONCLUSION

In view of the foregoing, Applicants believe all claims now pending in this Application are in condition for allowance. The issuance of a formal Notice of Allowance at an early date is respectfully requested.

Applicants believe that no fee is required. However, if a fee is required, the Commissioner is authorized to deduct such fee from the undersigned's Deposit Account No. 20-1430. Please deduct any additional fees from or credit any overpayment to, the above-noted Deposit Account.

If a telephone conference would expedite prosecution of this application, the Examiner is invited to telephone the undersigned at 415-576-0200.

Respectfully submitted,

  
Siegfried Ruppert  
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Attachments (Exhibit A; Rule 1.131 Declaration, including Exhibits 1-11)  
SIR:lo  
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## PROVISIONAL PATENT APPLICATION

In the name of the inventor

Weizhen Ji

**832C. Novel Splice Variant of Dishevelled-3-like Proteins  
and Nucleic Acids Encoding Same**



## Novel Splice Variant of Dishevelled-3-like Proteins and Nucleic Acids Encoding Same

The present invention discloses a novel protein encoded by a cDNA and/or by genomic DNA and proteins similar to it, namely, new proteins bearing sequence similarity to Dishevelled-3, nucleic acids that encode these proteins or fragments thereof, and antibodies that bind immunospecifically to a protein of the invention.

### Background

The *Drosophila* dishevelled gene (*dsh*) encodes a cytoplasmic phosphoprotein (Klingensmith et al., 1994) that regulates cell proliferation, acting as a transducer molecule for developmental processes, including segmentation and neuroblast specification. Pizzuti et al. (1996) noted that *dsh* is required for the function of the wingless gene product *wg*, a segment polarity gene homologous to the mammalian protooncogene *WNT1* (164820). The Dishevelled specific domain, specific to the signaling protein disheveled, is found adjacent to the PDZ domain (IPR001478), often in conjunction with DEP (IPR000591) and DIX (IPR001158). Pizzuti et al. (1996) reported the isolation and chromosomal mapping of 2 human *dsh* homologs, designated DVL1 and DVL3 by them. The human *dsh* homologs were isolated from a fetal brain cDNA library. DVL3 encodes a predicted 716-amino acid polypeptide that shows 74% nucleotide homology with human DVL1 and 71% homology with the mouse *Dvl1* gene. DVL1 and DVL3 share 64% amino acid identity. Pizzuti et al. (1996) reported that homology is particularly high in the N-terminal region and that there is more divergence in the C-terminal regions. PCR carried out using DNA from rodent human somatic cell hybrids and DVL3 specific primers led to the assignment of DVL3 to human chromosome 3. Pizzuti et al. (1996) regionally assigned DVL3 to band 3q27 using fluorescence in situ hybridization. Hybridization of poly(A) mRNA with the DVL3 cDNA revealed a 2.9-kb transcript with abundant expression in skeletal muscle, pancreas and heart. They also detected 5.9-kb and 5.0-kb transcripts in skeletal muscle, adult liver, adult heart, pancreas, and placenta. The 5.9-kb form was abundant in fetal tissues but the 5.0-kb form was absent from these tissues. Pizzuti et al. (1996) noted that Charcot-Marie-Tooth type 2B maps to chromosome 3q.

Bui et al. (1997) also isolated human DVL3, which shares 98% amino acid identity with mouse *Dvl3* and 49% with *Drosophila* *dsh*. The authors confirmed the chromosomal localization at 3p27. Semenov and Snyder (1997) isolated 3 human genes encoding proteins homologous to *Drosophila* *dsh*. The cDNA sequence of DVL3 reported by Semenov and Snyder (1997) differs from the previously reported sequences deposited in GenBank. Bui et al. (1997) detected expression of DVL3 mRNA in B cells, breast, kidney, bladder, endometrium, and 2 primary endometrial cultures. It was detected equally in normal human breast tissues and tumors and in colorectal samples of normal tissues, polyps, and tumors.

The sequence disclosed in the application represents a splice variant of human dishevelled 3 (DVL3), lacking a 363 bp long coding region containing a PDZ domain.

### References

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2. Klingensmith, J.; Nusse, R.; Perrimon, N. : The *Drosophila* segment polarity gene dishevelled encodes a novel protein required for response to wingless signal. *Genes Dev.* 8: 118-130, 1994. PubMed ID : 8288125

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### Brief Description of the Drawings

Figure 1. Nucleotide sequence encoding the Dishevelled-3-like protein (Acc. No. CG164330-01) of the invention.

Figure 2. Protein sequence encoded by the nucleotide sequence shown in Figure 1.

Figure 3A. A high-scoring match as determined by a BLASTN search of GenBank Composite (no HTG) dated 12/21/01 using the sequence of the Dishevelled-3-like gene of the invention.

Figure 3B. A high-scoring match as determined by a BLASTP search (versus Non-Redundant Composite dated 12/21/01) using the sequence of the Dishevelled-3-like protein of the invention.

Figure 3C. BLASTN identity search of CuraGen Corporation's human SeqCalling database using the Dishevelled-3-like gene of the invention.

Figure 4. ClustalW alignment of the protein of Acc. No. CG164330-01 with similar Dishevelled-3s.

Figure 5: PSORT, SignalP and hydropathy results for the Dishevelled-3-like protein of Acc. No. CG164330-01.

### Description of the Invention

#### Method of Identifying the Nucleic Acid Encoding the Dishevelled-3-Like Protein.

The sequence of Acc. No. CG164330-01 was derived by laboratory cloning of cDNA fragments, by *in silico* prediction of the sequence. cDNA fragments covering either the full length of the DNA sequence, or part of the sequence, or both, were cloned. *In silico* prediction was based on

sequences available in CuraGen's proprietary sequence databases or in the public human sequence databases, and provided either the full length DNA sequence, or some portion thereof.

The laboratory cloning was performed using one or more of the methods summarized below:

SeqCalling™ Technology: cDNA was derived from various human samples representing multiple tissue types, normal and diseased states, physiological states, and developmental states from different donors. Samples were obtained as whole tissue, primary cells or tissue cultured primary cells or cell lines. Cells and cell lines may have been treated with biological or chemical agents that regulate gene expression, for example, growth factors, chemokines or steroids. The cDNA thus derived was then sequenced using CuraGen's proprietary SeqCalling technology. Sequence traces were evaluated manually and edited for corrections if appropriate. cDNA sequences from all samples were assembled together, sometimes including public human sequences, using bioinformatic programs to produce a consensus sequence for each assembly. Each assembly is included in CuraGen Corporation's database. Sequences were included as components for assembly when the extent of identity with another component was at least 95% over 50 bp. Each assembly represents a gene or portion thereof and includes information on variants, such as splice forms, single nucleotide polymorphisms (SNPs), insertions, deletions and other sequence variations.

Variant sequences are also included in this application. A variant sequence can include a single nucleotide polymorphism (SNP). A SNP can, in some instances, be referred to as a "cSNP" to denote that the nucleotide sequence containing the SNP originates as a cDNA. A SNP can arise in several ways. For example, a SNP may be due to a substitution of one nucleotide for another at the polymorphic site. Such a substitution can be either a transition or a transversion. A SNP can also arise from a deletion of a nucleotide or an insertion of a nucleotide, relative to a reference allele. In this case, the polymorphic site is a site at which one allele bears a gap with respect to a particular nucleotide in another allele. SNPs occurring within genes may result in an alteration of the amino acid encoded by the gene at the position of the SNP. Intragenic SNPs may also be silent, when a codon including a SNP encodes the same amino acid as a result of the redundancy of the genetic code. SNPs occurring outside the region of a gene, or in an intron within a gene, do not result in changes in any amino acid sequence of a protein but may result in altered regulation of the expression pattern. Examples include alteration in temporal expression, physiological response regulation, cell type expression regulation, intensity of expression, and stability of transcribed message.

One or more genomic clones AC048331, AC061705, AC092931 on chromosome 3 were identified by TBLASTN using CuraGen Corporation's sequence file for members of Dishevelled-3 and/or the Dishevelled family, run against the genomic daily files made available by GenBank or obtained from Human Genome Project Sequencing centers. These sequences were analyzed for putative coding regions as well as for similarity to known DNA and protein sequences. Programs used for these analyses include Grail, Genscan, BLAST, HMMER, FASTA, Hybrid and other relevant programs. Putative coding regions were spliced from the genomic clone and then concatenated using a known homolog for reference. The derived sequence may have been further extended using additional genomic clones showing greater than 98% identity to the open reading frame.

The regions defined by the procedures described above were then manually integrated and corrected for apparent inconsistencies that may have arisen, for example, from misaligned bases in the original fragments or from discrepancies between predicted exon junctions, and regions of sequence similarity, to derive the final sequence disclosed herein. When necessary, the process to identify and analyze genomic clones was reiterated to derive the full length sequence. The following public components were thus included in the invention: AC048331, AC061705, AC092931.

The DNA sequence was analyzed to identify any open reading frames encoding novel full length proteins as well as novel splice forms of these genes. The DNA sequence and protein sequence for a novel Dishevelled-3-like gene are reported here as CuraGen Acc. No. CG164330-01.

## Results

The novel nucleic acid of 2634 nucleotides (designated CuraGen Acc. No. CG164330-01) encoding a novel Dishevelled-3-like protein is shown in Fig. 1. An open reading frame was identified beginning at nucleotides 51-53 and ending at nucleotides 1836-1838. This open reading frame begins with an ATG initiation codon and ends with a TGA stop codon. This polypeptide represents a novel functional Dishevelled-3-like protein. The start and stop codons of the open reading frame are highlighted in bold type. Putative untranslated regions (underlined), if any, are found upstream from the initiation codon and downstream from the termination codon. The encoded protein having 595 amino acid residues is presented using the one-letter code in Fig. 2.

## Similarities

In a search of sequence databases, it was found, for example, that the nucleic acid sequence of this invention has 1325 of 1501 bases (88%) identical to a gb:GENBANK-ID:AF006013|acc:AF006013.1 mRNA from Homo sapiens (Homo sapiens dishevelled 3 (DVL3) mRNA, complete cds) (Fig. 3A). The full amino acid sequence of the protein of the invention was found to have 336 of 336 amino acid residues (100%) identical to, and 336 of 336 amino acid residues (100%) similar to, the 716 amino acid residue ptnr:SWISSPROT-ACC:Q92997 protein from Homo sapiens (Human) (Segment polarity protein dishevelled homolog DVL-3 (Dishevelled-3) (DSH homolog 3))(Fig. 3B).

A multiple sequence alignment is given in Fig. 4, with the protein of the invention being shown on the first line in a ClustalW analysis comparing the protein of the invention with related protein sequences. Please note this sequence represents a splice form of Dishevelled-3 as indicated in positions 260 to 381 aa.

The presence of identifiable domains in the protein disclosed herein was determined by searches versus domain databases such as Pfam, PROSITE, ProDom, Blocks or Prints and then identified by the Interpro domain accession number. Significant domains are summarized in Table 1.

Scores for sequence family classification (score includes all domains):

Model	Description	Score	E-value	N
<u>DIX</u> ( <u>InterPro</u> )	DIX domain	194.5	1.7e-54	1
<u>Dishevelled</u> ( <u>InterPro</u> )	Dishevelled specific domain	136.6	4.5e-37	1
<u>DEP</u> ( <u>InterPro</u> )	Domain found in Dishevelled, Egl-10, and	121.1	2e-32	1
<u>oxidored_g1</u> ( <u>InterPro</u> )	NADH-Ubiquinone/plastoquinone (complex I)	3.5	5.1	1

## Parsed for domains:

Model	Domain	seq-f	seq-t	hmm-f	hmm-t	score	E-value
DIX	1/1	1	82	1	86	194.5	1.7e-54
Dishevelled	1/1	142	213	1	74	136.6	4.5e-37
oxidored_g1	1/1	245	272	291	316	3.5	5.1
DEP	1/1	301	375	1	89	121.1	2e-32

## describe domains and functional relevance

Dishevelled (Dsh) protein is an important component of the Wnt signal-transduction pathway. It has three relatively conserved domains: DIX, PDZ and DEP. The DIX domain of Dvl-1 (a mammalian Dishevelled homolog) shares 37% identity with the C-terminal region of Axin. Dsh can interact with the Axin/APC/GSK3/beta-catenin complex, and may thus modulate its activity.

The Wnt signaling pathway is conserved in various species from worms to mammals, and plays important roles in development, cellular proliferation, and differentiation. The molecular mechanisms by which the Wnt signal regulates cellular functions are becoming increasingly well understood. Wnt stabilizes cytoplasmic beta-catenin, which stimulates the expression of genes including c-myc, c-jun, fra-1, and cyclin D1. Axin and its homolog Axil are components of the Wnt signaling pathway that negatively regulate this pathway. Other components of the Wnt signaling pathway, including Dvl, glycogen synthase kinase-3beta (GSK-3beta), beta-catenin, and adenomatous polyposis coli (APC), interact with Axin, and the phosphorylation and stability of beta-catenin are regulated in the Axin complex. Axil has similar functions to Axin. Thus, Axin and Axil act as scaffold proteins in the Wnt signaling pathway, thereby modulating the Wnt-dependent cellular functions.

The Dishevelled specific domain is specific to the signaling protein dishevelled. In *Drosophila*, the dishevelled segment polarity protein is required to establish coherent arrays of polarized cells and segments in embryos. It plays a role in wingless signaling, possibly through the reception of the wingless signal by target cells and subsequent redistribution of arm protein in response to that signal in embryos. The domain is found adjacent to the PDZ domain (IPR001478), often in conjunction with DEP (IPR000591) and DIX (IPR001158).

This indicates that the sequence of the invention has properties similar to those of other proteins known to contain this/these domain(s) and similar to the properties of these domains.

### Chromosomal information:

The Dishevelled-3-like gene disclosed in this invention maps to chromosome 3. This assignment was made using mapping information associated with genomic clones, public genes and ESTs sharing sequence identity with the disclosed sequence and CuraGen Corporation's Electronic Northern bioinformatic tool.

### Tissue expression

The Dishevelled-3-like gene disclosed in this invention is expressed in at least the following tissues: fetal brain, fetal liver/spleen, melanocyte, placenta, ovary (tumor), breast, fetal heart, colon, uterus (pregnant), brain-hippocampus, embryo, parathyroid gland (tumor), heart, fetal lung. Expression information was derived from the tissue sources of the sequences that were included in the derivation of the sequence of CuraGen Acc. No. CG164330-01.

### Cellular Localization and Sorting

The PSORT, SignalP and hydropathy profile for the Dishevelled-3-like protein are shown in Fig. 5. The results predict that this sequence has no signal peptide and is likely to be localized in the nucleus with a certainty of 0.7000 predicted by PSORT. The hydropathy profile is characteristic of this gene family.

### Functional Variants and Homologs

The novel nucleic acid of the invention encoding a Dishevelled-3-like protein includes the nucleic acid whose sequence is provided in Fig. 1, or a fragment thereof. The invention also includes a mutant or variant nucleic acid any of whose bases may be changed from the corresponding base shown in Fig. 1 while still encoding a protein that maintains its Dishevelled-3-like activities and physiological functions, or a fragment of such a nucleic acid. The invention further includes nucleic acids whose sequences are complementary to the sequence of CuraGen Acc. No. CG164330-01, including nucleic acid fragments that are complementary to any of the nucleic acids just described. The invention additionally includes nucleic acids or nucleic acid fragments, or complements thereto, whose structures include chemical modifications. Such modifications include, by way of non-limiting example, modified bases, and nucleic acids whose sugar phosphate backbones are modified or derivatized. These modifications are carried out at least in part to enhance the chemical stability of the modified nucleic acid, such that they may be used, for example, as antisense binding nucleic acids in therapeutic applications in a subject. In the mutant or variant nucleic acids, and their complements, up to about 12% of the bases may be so changed.

The novel protein of the invention includes the Dishevelled-3-like protein whose sequence is provided in Fig. 2. The invention also includes a mutant or variant protein any of whose residues may be changed from the corresponding residue shown in Fig. 2 while still encoding a protein that maintains its Dishevelled-3-like activities and physiological functions, or a functional

fragment thereof. In the mutant or variant protein, up to about 0% of the amino acid residues may be so changed.

### Chimeric and Fusion Proteins

The present invention includes chimeric or fusion proteins of the Dishevelled-3-like protein, in which the Dishevelled-3-like protein of the present invention is joined to a second polypeptide or protein that is not substantially homologous to the present novel protein. The second polypeptide can be fused to either the amino-terminus or carboxyl-terminus of the present CG164330-01 polypeptide. In certain embodiments a third nonhomologous polypeptide or protein may also be fused to the novel Dishevelled-3-like protein such that the second nonhomologous polypeptide or protein is joined at the amino terminus, and the third nonhomologous polypeptide or protein is joined at the carboxyl terminus, of the CG164330-01 polypeptide. Examples of nonhomologous sequences that may be incorporated as either a second or third polypeptide or protein include glutathione S-transferase, a heterologous signal sequence fused at the amino terminus of the Dishevelled-3-like protein, an immunoglobulin sequence or domain, a serum protein or domain thereof (such as a serum albumin), an antigenic epitope, and a specificity motif such as (His)<sub>6</sub>.

The invention further includes nucleic acids encoding any of the chimeric or fusion proteins described in the preceding paragraph.

### Antibodies

The invention further encompasses antibodies and antibody fragments, such as Fab, (Fab)<sub>2</sub>, or single chain FV constructs, that bind immunospecifically to any of the proteins of the invention. Also encompassed within the invention are peptides and polypeptides comprising sequences having high binding affinity for any of the proteins of the invention, including such peptides and polypeptides that are fused to any carrier particle (or biologically expressed on the surface of a carrier) such as a bacteriophage particle.

### Uses of the Compositions of the Invention

The protein similarity information, expression pattern, cellular localization, and map location for the protein and nucleic acid disclosed herein suggest that this Dishevelled-3-like protein may have important structural and/or physiological functions characteristic of the Dishevelled family. Therefore, the nucleic acids and proteins of the invention are useful in potential diagnostic and therapeutic applications and as a research tool. These include serving as a specific or selective nucleic acid or protein diagnostic and/or prognostic marker, wherein the presence or amount of the nucleic acid or the protein are to be assessed. These also include potential therapeutic applications such as the following: (i) a protein therapeutic, (ii) a small molecule drug target, (iii) an antibody target (therapeutic, diagnostic, drug targeting/cytotoxic antibody), (iv) a nucleic acid useful in gene therapy (gene delivery/gene ablation), (v) an agent promoting tissue regeneration *in vitro* and *in vivo*, and (vi) a biological defense weapon.

The nucleic acids and proteins of the invention have applications in the diagnosis and/or treatment of various diseases and disorders. For example, the compositions of the present



invention will have efficacy for the treatment of patients suffering from: adrenoleukodystrophy, Alzheimer's disease, autoimmune disease, allergies, addiction, anxiety, ataxia-telangiectasia, asthma, ARDS, atherosclerosis, behavioral disorders, aortic stenosis, atrial septal defect (ASD), atrioventricular (A-V) canal defect, ductus arteriosus, allergy, cerebral palsy, congenital adrenal hyperplasia, cirrhosis, cardiomyopathy, congenital heart defects, diabetes, diverticular disease, epilepsy, emphysema, endometriosis, endocrine dysfunctions, graft versus host disease, glomerulonephritis, graft versus host disease (GVHD), growth and reproductive disorders, hemophilia, hypercoagulation, hypercalcemia, Huntington's disease, hypertension, hypogonadism, fertility, idiopathic thrombocytopenic purpura, immunodeficiencies, interstitial nephritis, IgA nephropathy, lymphoma, inflammatory bowel disease, Lesch-Nyhan syndrome, leukodystrophies, multiple sclerosis, muscular dystrophy, myasthenia gravis, neurodegeneration, neuroprotection, obesity, Parkinson's disease, pain, polycystic kidney disease, pulmonary stenosis, pancreatitis, renal artery stenosis, renal tubular acidosis, stroke, systemic lupus erythematosus, scleroderma, subaortic stenosis, transplantation, tuberous sclerosis, Von Hippel-Lindau (VHL) syndrome, ventricular septal defect (VSD), valve diseases, Von Hippel-Lindau (VHL) syndrome, ulcers, cancers as well as other diseases, disorders and conditions.

These materials are further useful in the generation of antibodies that bind immunospecifically to the novel substances of the invention for use in diagnostic and/or therapeutic methods.

## FIGURES

Figure 1. Nucleotide sequence encoding the Dishevelled-3-like protein of the invention.

>CG164330\_01

GGCGCGCCGAGCAGGCGCGCGCGGGCGCGCGGCCGAGGCGACAGCCATGGGGGAGA	60
CCAAGATCATCTACCACTTGGATGGGCGAGGACAGCCGCTACTTGTGAAGCTGCCTCCG	120
CGCGCGGACGCGCTCACCCTTGTGGGAGCTTTAAGGCGCTTTCGCGACGCCAGCATTAAGT	180
TTCTCTTCAAGTCTATGAGGACAGGATTTGGAGTGTGGTAGAGGAGATCTAGTGAAGACA	240
ATGCCAAGCTACACCTGCTCAAAGTGGCGGGGTGGTGTCTGCGTGTGTGTCATCGGATGCT	300
CACACCGACCGACCGCCCTCTCTGTGTGTGCTTAACCTTCCTGGAGCTGCCACACATATGG	360
AGCGCACGGGAGGACATGGGGACATCCGACGCCCACTTCCACACCTCATCTGTGTGGTGGG	420
CGACGACGAGAGAAATCTGGACATATGCCACAGACAGACGGACTCTTTGGTGTCTGCCACAGG	480
GGCGGCCACGCGCGGAGGAGATGGCCAGAGCTCAACCCGAGTAAGTAAGGAATCGGAGG	540
GGAAACGGGCGGAGGACAGGGGGTATGATAGATCTATCCACCACTTATGACCACTGAGC	600
TGGACACCCAGCTCTTTGACTCAGATGAGGATGACTCCACCGCTTGCATCGACTCT	660
CCACAGAACAGAGCAGTGCCTCACGCTGTAGAAAGACACAAGCGCGGCGCGGAGC	720
AGATGGGTTCCTGGATGTAGCGCGTCTCGTCTCTTCAGCAGCATCACGGAATCCACCAT	780
CACTCAACATCTACCGGTCACTCTCAACATGGAAAATAATAACTTCTTGCACCATCA	840
CTTCCACAGCTCTCTCATACAGCTTCCATCTCTGACAGAGAGCGCTTAGACAGATTC	900
ACTTGTCTATCCACAGTACATGGTGTGCATCTGTAAAGCGCATGGCCTCGCTGAATAC	960
GTGTGGAGGTTCGCTGACGCATATGGCTCAAGATTAACATCTCTAATGCTTATCGGCT	1020
CAGATGTGTGGTACCTGTGCTTACCAATCTGTGGAGCTCTCACGACCGGAGGAGAGGCC	1080
GCAAGTCTGCGTACGACTCTCTGAAAGCTGGCTTCATCCGCGCATACCGCTCAACAAGTCA	1140
CTTCTTCGAGCAGGCTCTACTACATCTCGTGACCTCTCGCGCAACATGGCCACACTGT	1200
CTTCCACAGCTACGATGGCTCTGACGTGGGCGCTCTGACACGAGACAGTGGCCCTTTGC	1260
CGCCACCGGGGCGCCCTCTTGGCCATGGCTTTCGAGTACAGTACCCGCAACCCCGC	1320
ACCCATPACAAACCCGCTCGGCTTCTCCGAGCTGGGCTACAGCTACGGCGGGGCGAGG	1380
CCAGCATGTACGACAGCAGGAGCGAGTGTGAGACGCTGGCTCAACCTGTGGCGGACGAT	1440
GGAGAGAGGAGAAAGACCCAAGAGCGGGGACTCCAAGTCTGGGGGACGGCGCGAGAT	1500
CGGACGACACACACAGCAGAGCTCTGGGGGCGCGGGAGCGGGCGCGAGCGCT	1560
CAGGGCCGCGCCAGCGAGACAGCCACACCGGACCACTTCTCGGCACAGCAAGCTCT	1620
GCAGCTACACATACACCCGAGTACGGTCTCTCCGAGTGGCGCCCTCTTACGGCGCCC	1680
CATGCTGTATGTGCCCGCCGCGCCGCTGCTTGGGCGCCCGAGGACCGCTTACGGCC	1740

```

GCGACCTGGCCTCAGTGCCTCCGGAAGTACCGCCAGCAGACAGTCTCTCCGCATGGCCA 1800
TGGGAAACCCAGTGAGTTCTTTGTGGATGTAGTGTGAGCAGGCGCCCTCCCCAGCTCC 1860
ATTTCGCTCCACCCAGCCGCGCTGCGTTCCTCTCTCCATCCGTCCTCTTTTACTTT 1920
GTCTGGTACCTGAAAGGAAATAAAAGGAACATAATCCAGGTGCGCTAACTGCTCGCAGG 1980
GTGCTGCGAGGGTGGGGTGACCTACCGATTGGCTCTGACGCCCTTAACCTGCCCTCTGG 2040
CCCCAGTTCGTTTCCTCTGCCCACTAATCCCTGCGCAGGACTTCCAGGAGCCCTTTTGT 2100
CTCTGGGACCAGACTGTGTGGTGTACCCCTTACTCCCTCTGCAACCCCATTTTGGGA 2160
GTGTGACCCCAAGCAATGACCTTGGTGGCAGCTCACTCCCTCATTTCTCTGTTTCCCTTT 2220
AGCTCCCTTTACCAATTATTCAGCTACATCATCCCTCTATTAAACCCACCCCATCAGGC 2280
ACGTGTGCAAACTCTTGACTTTACCCACATTAAGTAAACCAAAATATTAITTTGCTTCAT 2340
CTGCCCTACTAACCATCCCTGCTGCTGCTCAGTCTGCAACCTAAAGCTGTAGTCT 2400
GCCTCAATAGCCATCCATGCCATCCCTGCCCTGTGCTAGATCAGAGGCCAGAGGGCCC 2460
CTCCAGTTGCCGTGACAGCTGGTGGCTTCCAGGAGGAGTCTGTGCTTACCCCTGCCCA 2520
CCTCTGCCCTGCGTGGTGTCTCTTCAGACCCCTAACCTACTAACAGCAGGCTCATCT 2580
CACCTTCAGGCTGAACATTTCTTTTCTTTCTTTTTCCTCCCCCAATTACCC 2634

```

**Figure 2. Protein sequence encoded by the nucleotide sequence shown in Figure 1.**

```

>CG164330.01
MGETKTIYHLDDQETFPYLVKLELPAERVTLADFVKVLQRPSTKFFFSKDDDFGVKKEI 60
SDDNAKLPCFNFRVSVWLVSAEGSHPDPAFFCADNPSELPPMERTGGIGDSRFPSPFPH 120
AGGGQENLNDNDETDSLVAQRGRFRRRDGFPAHRLNGTAKGERRRGGVYDSSSTLM 180
SSELETTSFDDDEDDSTRFSSSTBQSSASRLMRKRKRKRKRKRKRKRKRKRKRKRKRKR 240
STMSLNIIITVTNMEKYNFLSTTTSTSSSTTSSIPDTERLDLPHLSIHSMDMAIVKAMAS 300
PESGLEVRDRLWKITIPNFIGSDVVDWLYHNVEGFTDRREARKYASNLKAGVFIHRTV 360
NKITPSGCQCYIFGDLGNMANLSLHDHSGSGASDQDTLAPLPHPGAAPWPMAPFYQYP 420
PPPHYPNHPFGPELGYSGGSSASSQHSBGRSSSGSNRSGSDRRKEKDPKAGDSKSGGS 480
GSESDHTTRSSLRGPRERAPSERSGPAASEHSHRSHSLASSLSRHHTHPSYGPFGVPPL 540
YGPMLMPPPPAAMPFGAPPGRDLASVPPELTASRQSFMRMAGNPFSEFVDVM 595

```

**Figure 3A. BLASTN search using CuraGen Acc. No. CG164330.01.**

```

>gb:GENBANK-ID:AF006013|acc:AF006013.1 Homo sapiens dishevelled 3 (DVL3) mRNA,
complete cds - Homo sapiens, 2286 bp.
Length = 2286

```

**Plus Strand HSPs:**

Score = 5641 (846.4 bits), Expect = 4.8e-249, P = 4.8e-249

Identities = 1325/1501 (88%), Positives = 1325/1501 (88%), Strand = Plus / Plus

```

Query: 459 TCTTTGGTGCTGCCCCAGCGAGGCGCGCCAGCCG-GAGGGATGCCCCAGCAGCATGCAAC 517
TCTT GG TCT CC A G GGG C CG G G G TG C G GCAT C AC
Sbjct: 787 TCTTGGGACTCT-CC-ATTGTGGGCAAAGCAACGAGGTGTTGACGGCG-GCAT-CTAC 842

```

```

Query: 518 CC-GGCTAAATGGAAGTGCAGAGGGGGAACGGCGGC-GA-GGAC-CAGGGGGTTATGA-- 571
GGCT AT G G GGG GGC GC GA GGAC CA G G A GA
Sbjct: 843 ATTGGCTCTATCATGAAGGGT-GGGGCGTGCTGCTGATGACGCATCGAGCCAGGAGA 901

```

```

Query: 572 TA-GCTCATCCACCTTATGACGAGTGAGCTG-GAGACCACAGCTTCTTT-GACTCAGA 628
TA G T T CA A GAG A T A CT GAG A CA G G A T T GA CAG
Sbjct: 902 TATGTTGTGTACAGSTAAACGAG-A-TCAACTTTGAGAACATGAG-TAATGACGATGCAG- 957

```

```

Query: 629 TGAGGATGACTCCACAGCAGGTTTCAGCAGCTCCACAGAACAGAGCAGTGCCTCACGCC 688
T GG T ACT C AG GT CA CA CC G CA C TG CT GCC
Sbjct: 958 TCCGGG-ACTGCGGGAGATTGTGCA-CAAA-CCGGGGCCATCACCTGTAGTGTAGCA 1014

```

Query: 689 GATGAGAAGACACAAGCGGCGGGCGGGAAGCAGAAGTTTCTC-GGATTGAGCGG-TCC 746  
TG GAC CAAG C CG GG GC A TT C C GGA GAGC TCC

Sbjct: 1015 AGTGCTGGGACCAAGTC-CA-CGT-GGTTGCTTACATTTGCCAGGAGCGAGCCCATC 1071

Query: 747 TCGTCC-ITCAGCA-GCATCACGGA-CTCCACCA-TGTCACTCAACATCATCAGGTCAC 802  
G CC TT A C GC C GG CTCC CA TG CA A C CAC TC C

Sbjct: 1072 G-GCCCATTGACCCCTGGGCGCTGGGTCTCCCACTG-CAGCCATGACCGGACCATTTCC 1129

Query: 803 TCTCAACATGGAAAAATATAACTTCTTGAGCACCATCACCTCCACCAGCTCTCCATCAC 862  
T CA A GG A A CT C TGAGCACCATCACCTCCACCAGCTCTCCATCAC

Sbjct: 1130 TG-CAT-ACGGCATGAGCCC-CTCCTGAGCACCATCACCTCCACCAGCTCTCCATCAC 1186

Query: 863 CAGTTCATCCCTGACACAGAGCGCCTAGACGACTTCCACTTGTCCATCCACAGTGACAT 922  
CAGTTCATCCCTGACACAGAGCGCCTAGACGACTTCCACTTGTCCATCCACAGTGACAT

Sbjct: 1187 CAGTTCATCCCTGACACAGAGCGCCTAGACGACTTCCACTTGTCCATCCACAGTGACAT 1246

Query: 923 GGCTGCCATCGTAAAAGCCATGGCCTCCCTGAATCAGGGTTGGAGGTCGTCAGCCGAT 982  
GGCTGCCATCGTAAAAGCCATGGCCTCCCTGAATCAGGGTTGGAGGTCGTCAGCCGAT

Sbjct: 1247 GGCTGCCATCGTAAAAGCCATGGCCTCCCTGAATCAGGGTTGGAGGTCGTCAGCCGAT 1306

Query: 983 GTGGCTCAAGATTACCATCCCTAATGCTTTTCATCGGCTCAGATGTGGTGGACTGGCTGTA 1042  
GTGGCTCAAGATTACCATCCCTAATGCTTTTCATCGGCTCAGATGTGGTGGACTGGCTGTA

Sbjct: 1307 GTGGCTCAAGATTACCATCCCTAATGCTTTTCATCGGCTCAGATGTGGTGGACTGGCTGTA 1366

Query: 1043 CCACAATGTGGAAGGCTTTCACGGACCGGAGGAGGCCCGCAAGTATGCCAGCAACCTGTCT 1102  
CCACAATGTGGAAGGCTTTCACGGACCGGAGGAGGCCCGCAAGTATGCCAGCAACCTGTCT

Sbjct: 1367 CCACAATGTGGAAGGCTTTCACGGACCGGAGGAGGCCCGCAAGTATGCCAGCAACCTGTCT 1426

Query: 1103 GAAAGCTGGCTTCATCCGCCATACCGTCAACAAGATCACCTTCTCCGAGCAGTGCTACTA 1162  
GAAAGCTGGCTTCATCCGCCATACCGTCAACAAGATCACCTTCTCCGAGCAGTGCTACTA

Sbjct: 1427 GAAAGCTGGCTTCATCCGCCATACCGTCAACAAGATCACCTTCTCCGAGCAGTGCTACTA 1486

Query: 1163 CATCTTCGGTGACCTCTGCGGCAACATGGCCAACTGTCTCTCCAGATCAAGATGGCTC 1222  
CATCTTCGGTGACCTCTGCGGCAACATGGCCAACTGTCTCTCCAGATCAAGATGGCTC

Sbjct: 1487 CATCTTCGGTGACCTCTGCGGCAACATGGCCAACTGTCTCTCCAGATCAAGATGGCTC 1546

Query: 1223 CAGTGGCGCCTCTGACAGGACACACTGGCCCTTTTGCCGACCCGGGGGCGCCCTTG 1282  
CAGTGGCGCCTCTGACAGGACACACTGGCCCTTTTGCCGACCCGGGGGCGCCCTTG

Sbjct: 1547 CAGTGGCGCCTCTGACAGGACACACTGGCCCTTTTGCCGACCCGGGGGCGCCCTTG 1606

Query: 1283 GCCCATGGCTTTCCCGTACCAGTACCCGCCACCCCGCACCCATACAACCGCACCCGGG 1342  
GCCCATGGCTTTCCCGTACCAGTACCCGCCACCCCGCACCCATACAACCGCACCCGGG

Sbjct: 1607 GCCCATGGCTTTCCCGTACCAGTACCCGCCACCCCGCACCCATACAACCGCACCCGGG 1666

Query: 1343 CTTCCCGGAGCTGGGCTACAGTACGGCGGGGCGAGCGCCAGCAGTCAGCACAGCGAAG 1402  
CTT GGAGCTGGGCTACAGTACGGCGGGGCGAGCGCCAGCAGTCAGCACAGCGAAG

Sbjct: 1667 CTTGGGGGAGCTGGGCTACAGTACGGCGGGGCGAGCGCCAGCAGTCAGCACAGCGAAG 1726

Query: 1403 CAGTCGGAGCAGTGCTCTCAACCGTAGCGGCAGCAGTCGGAGGAAGGAGAAGGACCCGAA 1462  
CAGTCGGAGCAGTGCTCTCAACCGTAGCGGCAGCAGTCGGAGGAAGGAGAAGGACCCGAA

Sbjct: 1727 CAGTCGGAGCAGTGCTCTCAACCGTAGCGGCAGCAGTCGGAGGAAGGAGAAGGACCCGAA 1786

Query: 1463 GGC CGGGGACTCCAAGTCCGGGGGCGAGCGGCAGCGAATCGGACCACACCAACGAGCAG 1522  
GGC CGGGGACTCCAAGTCCGGGGGCGAGCGGCAGCGAATCGGACCACACCAACGAGCAG

Sbjct: 1787 GGC CGGGGACTCCAAGTCCGGGGGCGAGCGGCAGCGAATCGGACCACACCAACGAGCAG 1846

Query: 1523 CCGCGGGGCGCGGGGAGCGGGCGCCAGCGAGCGCTCAGGGCGCGGGCCAGCGAGCA 1582

[illegible]

```

Query:   400 TCCACCCCTCATGCTGGTGGGGGCGCCAGGAGAACTGGACAAATGACACAGAGACGGACT 459
          |||
Sbjct:   361 TCCACCCCTCATGCTGGTGGGGGCGCCAGGAGAACTGGACAAATGACACAGAGACGGACT 420

Query:   460 CTTTGGTGCTGCCACGCGAGGGCGGCCACGCCGGAGGGATGGCCCAGAGCATGCAACCC 519
          |||
Sbjct:   421 CTTTGGTGCTGCCACGCGAGAGCGGCCACGCCGGAGGGATGGCCCAGAGCATGCAACCC 480

Query:   520 GGCTAAATGGAAGTGCAGAGGGGGAACGCGCGCGAGGACCAGGGGGTTATGATAGCTCAT 579
          |||
Sbjct:   481 GGCTAAATGGAAGTGCAGAGGGGGAACGCGCGCGAGAACAGGGGGTTATGATAGCTCAT 540

Query:   580 CCACCCCTTATGAGCAGTGAGCTGGAGACCAACAGCTTCTTTGACTCAGATGAGGATGACT 639
          |||
Sbjct:   541 CCACCCCTTATGAGCAGTGAGCTGGAGACCAACAGCTTCTTTGACTCAGATGAGGATGACT 600

Query:   640 CCACCAGCAGGTTTCAGCAGCTCCACAGAACAGAGCAGTGCCCTCACGCTGATGAGAAGAC 699
          |||
Sbjct:   601 CCACCAGCAGGTTTCAGCAGCTCCACAGAACAGAGCAGTGCCCTCACGCTGATGAGAAGAC 660

Query:   700 ACAAGCGCGCGGCGCGGAAGCAGAGGTTTCTCGGATTGAGCGGCTCTCGTCTCTCAGCA 759
          |||
Sbjct:   661 ACAAGCGCGCGGCGCGGAAGCAGAGGTTTCTCGGATTGAGCGGCTCTCGTCTCTCAGCA 720

Query:   760 GCATCACGGACTCCACCATGTCTACTCAACATCATCACGGTCACTCTCAACATGGA AAAAT 819
          |||
Sbjct:   721 GCATCACGGACTCCACCATGTCTACTCAACATCATCACGGTCACTCTCAACATGGA AAAAT 780

Query:   820 ATAACCTCTTGAGCACCATCA 840
          |||
Sbjct:   781 ATAACCTCTTGAGCATCTCCA 801

```

**Figure 3B. BLASTP search using the protein of CuraGen Acc. No. CG164330-01.**

```

>ptnr:SWISSPROT-ACC:Q92997 Segment polarity protein dishevelled homolog DVL-3
      (Dishevelled-3) (DSH homolog 3) - Homo sapiens (Human), 716 aa.
      Length = 716

```

Score = 1811 (637.5 bits), Expect = 0.0, Sum P(2) = 0.0  
 Identities = 336/336. (100%), Positives = 336/336 (100%)

```

Query:   260 LSTITSTSSSITSSIPDTERLDDFHLSDMAAIVKAMASPESGLEVRDRMLKITIPN 319
          LSTITSTSSSITSSIPDTERLDDFHLSDMAAIVKAMASPESGLEVRDRMLKITIPN
Sbjct:   381 LSTITSTSSSITSSIPDTERLDDFHLSDMAAIVKAMASPESGLEVRDRMLKITIPN 440

Query:   320 AFIGSDVVDWLYHNVEGFTDRREARKYASNLLKAGFIRHTVFNKITFSBQCYIIFGDLGCGN 379
          AFIGSDVVDWLYHNVEGFTDRREARKYASNLLKAGFIRHTVFNKITFSBQCYIIFGDLGCGN
Sbjct:   441 AFIGSDVVDWLYHNVEGFTDRREARKYASNLLKAGFIRHTVFNKITFSBQCYIIFGDLGCGN 500

Query:   380 MANLSLHDHDSGSGASDQDTLAPLPHGAAPWPMAPFYQYPPPPHYPNHPGFPPELGYSY 439
          MANLSLHDHDSGSGASDQDTLAPLPHGAAPWPMAPFYQYPPPPHYPNHPGFPPELGYSY
Sbjct:   501 MANLSLHDHDSGSGASDQDTLAPLPHGAAPWPMAPFYQYPPPPHYPNHPGFPPELGYSY 560

Query:   440 GGGASASSQHSEGRSSGNSRSGSDRRKEKDPKAGDSKSGSGSES DHTTRSSLRGPRERA 499
          GGGASASSQHSEGRSSGNSRSGSDRRKEKDPKAGDSKSGSGSES DHTTRSSLRGPRERA
Sbjct:   561 GGGASASSQHSEGRSSGNSRSGSDRRKEKDPKAGDSKSGSGSES DHTTRSSLRGPRERA 620

```

Query: 500 PSERSGPAASEHSHRSHSLASSLSRHHTHPSYGPPGVPLYGPPMLMPPPPAAMGPGP 559  
 PSERSGPAASEHSHRSHSLASSLSRHHTHPSYGPPGVPLYGPPMLMPPPPAAMGPGP  
 Sbjct: 621 PSERSGPAASEHSHRSHSLASSLSRHHTHPSYGPPGVPLYGPPMLMPPPPAAMGPGP 680

Query: 560 APPGRDLASVPPELTASRQSFRRMAMGNPSEFFVDVM 595  
 APPGRDLASVPPELTASRQSFRRMAMGNPSEFFVDVM  
 Sbjct: 681 APPGRDLASVPPELTASRQSFRRMAMGNPSEFFVDVM 716

Score = 1340 (471.7 bits), Expect = 0.0, Sum P(2) = 0.0  
 Identities = 258/260 (99%), Positives = 258/260 (99%)

Query: 1 MGETKIIYHLDGQETPYLVKLPLPAERVTLADFKGVLQRPYSYKFFFKSMDDDFGVVKEEI 60  
 MGETKIIYHLDGQETPYLVKLPLPAERVTLADFKGVLQRPYSYKFFFKSMDDDFGVVKEEI  
 Sbjct: 1 MGETKIIYHLDGQETPYLVKLPLPAERVTLADFKGVLQRPYSYKFFFKSMDDDFGVVKEEI 60

Query: 61 SDDNAKLPCFNGRVVSWLVSABGSHDPDAPPCADNPSELPPPMERTGGIGDSRPPSFH 120  
 SDDNAKLPCFNGRVVSWLVSABGSHDPDAPPCADNPSELPPPMERTGGIGDSRPPSFH  
 Sbjct: 61 SDDNAKLPCFNGRVVSWLVSABGSHDPDAPPCADNPSELPPPMERTGGIGDSRPPSFH 120

Query: 121 AGGGSQENLNDTETDSLVAQGRPRRRDGPHEATRLNGTAGGERRRPGGYDSSSTLM 180  
 AGGGSQENLNDTETDSLVAQGRPRRRDGPHEATRLNGTAGGERRRPGGYDSSSTLM  
 Sbjct: 121 AGGGSQENLNDTETDSLVAQGRPRRRDGPHEATRLNGTAGGERRRPGGYDSSSTLM 180

Query: 181 SSELETTTSFFDSDDEDSTSRFSSSTEQSSASRLMRHKKRRRRKQKVSRIERSSSFSSITD 240  
 SSELETTTSFFDSDDEDSTSRFSSSTEQSSASRLMRHKKRRRRKQKVSRIERSSSFSSITD  
 Sbjct: 181 SSELETTTSFFDSDDEDSTSRFSSSTEQSSASRLMRHKKRRRRKQKVSRIERSSSFSSITD 240

Query: 241 STMSLNIITVTILNMEKYNFL 260  
 STMSLNIITVTILNMEKYNFL  
 Sbjct: 241 STMSLNIITVTILNMEKYNFL 260

**Figure 3C. BLASTN identity search of CuraGen Corporation's Human SeqCalling database using CuraGen Acc. No. CG164330-01.**

>s3aq:239634112, 5183 bp.  
 Length = 5183

Minus Strand HSPs:

Score = 9052 (1358.2 bits), Expect = 0.0, Sum P(2) = 0.0  
 Identities = 2004/2176 (92%), Positives = 2004/2176 (92%), Strand = Minus / Plus

Query: 2634 GGTAATTTGGGGGAGGAAAAAAGAAAGAAAGAAATGTTTCAGGCCTGGAGGTGAGATGA 2575  
 GGTAATTTGGGGGAGGAAAAAAGAAAGAAAGAAATGTTTCAGGCCTGGAGGTGAGATGA  
 Sbjct: 2083 GGTAATTTGGGGGAGGAAAAAAGAAAGAAAGAAATGTTTCAGGCCTGGAGGTGAGATGA 2142

Query: 2574 GCCTGCTGGTTAGTAGGGTTAGGGGCTCTGAAGGAACACGACACGAGGCAGGCATGGGGC 2515  
 GCCTGCTGGTTAGTAGGGTTAGGGGCTCTGAAGGAACACGACACGAGGCAGGCATGGGGC  
 Sbjct: 2143 GCCTGCTGGTTAGTAGGGTTAGGGGCTCTGAAGGAACACGACACGAGGCAGGCATGGGGC 2202

Query: 2514 AGGGGTAGAGCAGAGATGCTCCCTGGAAGCCACACAGCTGCTCAGGCAACTGAGGGGGGCC 2455  
 AGGGGTAGAGCAGAGATGCTCCCTGGAAGCCACACAGCTGCTCAGGCAACTGAGGGGGGCC  
 Sbjct: 2203 AGGGGTAGAGCAGAGATGCTCCCTGGAAGCCACACAGCTGCTCAGGCAACTGAGGGGGGCC 2262

Query: 2454 TCTGGGCTCTGATCTAGGCACAGGCAGGCAGGCATGGCATGGATGGCTATTGGAGGCAGCTAC 2395  
 TCTGGGCTCTGATCTAGGCACAGGCAGGCAGGCATGGCATGGATGGCTATTGGAGGCAGCTAC

Sbjct: 2263 TCTGGGCTCTGATCTAGGCACAGGCAGGGATGGCATGGATGGCTATTGGAGGCGACTAC 2322

Query: 2394 AGCTTTAGGTTGCGAGACTGAGGCAGCAGGCAGGGGATGGTTAGTAGGGGCGAGATGAAG 2335  
AGCTTTAGGTTGCGAGACTGAGGCAGCAGGCAGGGGATGGTTAGTAGGGGCGAGATGAAG

Sbjct: 2323 AGCTTTAGGTTGCGAGACTGAGGCAGCAGGCAGGGGATGGTTAGTAGGGGCGAGATGAAG 2382

Query: 2334 CAAATATATTTTGGTTTCAGTAATGTGGGGTAAAGTCAAGAGGTTTGACACAGTGCCTGA 2275  
CAAATATATTTTGGTTTCAGTAATGTGGGGTAAAGTCAAGAGGTTTGACACAGTGCCTGA

Sbjct: 2383 CAAATATATTTTGGTTTCAGTAATGTGGGGTAAAGTCAAGAGGTTTGACACAGTGCCTGA 2442

Query: 2274 TGGGGTGGGTTAATAGAGGGATGATGTAGCTGAATAAATGGTGAAAGGGAGCTAAAGGG 2215  
TGGGGTGGGTTAATAGAGGGATGATGTAGCTGAATAAATGGTGAAAGGGAGCTAAAGGG

Sbjct: 2443 TGGGGTGGGTTAATAGAGGGATGATGTAGCTGAATAAATGGTGAAAGGGAGCTAAAGGG 2502

Query: 2214 GAAACGAGAGAATGAGGGAGTGAGCGTGCCACCAAGGTCATTGCTGGGGTCAACTCCCAA 2155  
GAAACGAGAGAATGAGGGAGTGAGCGTGCCACCAAGGTCATTGCTGGGGTCAACTCCCAA

Sbjct: 2503 GAAACGAGAGAATGAGGGAGTGAGCGTGCCACCAAGGTCATTGCTGGGGTCAACTCCCAA 2562

Query: 2154 AATGGGGGTTGCGAGAGGGAGTAAGGGGTAGCACCAACAAGTCTGGTCCAGAGACAAAA 2095  
AATGGGGGTTGCGAGAGGGAGTAAGGGGTAGCACCAACAAGTCTGGTCCAGAGACAAAA

Sbjct: 2563 AATGGGGGTTGCGAGAGGGAGTAAGGGGTAGCACCAACAAGTCTGGTCCAGAGACAAAA 2622

Query: 2094 GGGGTCCTGGGAAGTCTCGCGAGGGATTAGTGGGCAGAGAAACGAACCTGGGGCCAGAG 2035  
GGGGTCCTGGGAAGTCTCGCGAGGGATTAGTGGGCAGAGAAACGAACCTGGGGCCAGAG

Sbjct: 2623 GGGGTCCTGGGAAGTCTCGCGAGGGATTAGTGGGCAGAGAAACGAACCTGGGGCCAGAG 2682

Query: 2034 GCAGGTTAGGGGGCTGCGAGAGCCAATCGGTAGGTGCACCCACCCCTCGCAGCACCTGGG 1975  
GCAGGTTAGGGGGCTGCGAGAGCCAATCGGTAGGTGCACCCACCCCTCGCAGCACCTGGG

Sbjct: 2683 GCAGGTTAGGGGGCTGCGAGAGCCAATCGGTAGGTGCACCCACCCCTCGCAGCACCTGGG 2742

Query: 1974 AGCAGTTAGCGCACTGGATTAGTTCCCTTTTATTTCCCTTTTCAGGTACCGAGACAAAGTA 1915  
AGCAGTTAGCGCACTGGATTAGTTCCCTTTTATTTCCCTTTTCAGGTACCGAGACAAAGTA

Sbjct: 2743 AGCAGTTAGCGCACTGGATTAGTTCCCTTTTATTTCCCTTTTCAGGTACCGAGACAAAGTA 2802

Query: 1914 AAAAAGACGAGCGGATGGAGAGAGGAACCGAGCCGGCTGGGGTGGGAGCGGAATGGAGCT 1855  
AAAAAGACGAGCGGATGGAGAGAGGAACCGAGCCGGCTGGGGTGGGAGCGGAATGGAGCT

Sbjct: 2803 AAAAAGACGAGCGGATGGAGAGAGGAACCGAGCCGGCTGGGGTGGGAGCGGAATGGAGCT 2862

Query: 1854 GGGGAGGGGGCCTGCTCACATCACATCCACAAAGAATCACTGGGGTTTCCATGGCCA 1795  
GGGGAGGGGGCCTGCTCACATCACATCCACAAAGAATCACTGGGGTTTCCATGGCCA

Sbjct: 2863 GGGGAGGGGGCCTGCTCACATCACATCCACAAAGAATCACTGGGGTTTCCATGGCCA 2922

Query: 1794 TCGGAAGGACTGTCTGCTGGCGGTGAGTTCCGGGGCACTGAGGCCAGGTGCGGGCCCG 1735  
TCGGAAGGACTGTCTGCTGGCGGTGAGTTCCGGGGCACTGAGGCCAGGTGCGGGCCCG

Sbjct: 2923 TCGGAAGGACTGTCTGCTGGCGGTGAGTTCCGGGGCACTGAGGCCAGGTGCGGGCCCG 2982

Query: 1734 GAGGGGCTCCTGGGGGCCCATGAGCCGGCGGGCGGGGCAATCATGAGCTGGGGGGC 1675  
GAGGGGCTCCTGGGGGCCCATGAGCCGGCGGGCGGGGCAATCATGAGCTGGGGGGC

Sbjct: 2983 GAGGGGCTCCTGGGGGCCCATGAGCCGGCGGGCGGGGCAATCATGAGCTGGGGGGC 3042

Query: 1674 CGTAGAGAGGGGGCACTCCGGGAGGACCGTAGCTCGGGTGTGTGTGGTGGCTGCGAAGGC 1615  
CGTAGAGAGGGGGCACTCCGGGAGGACCGTAGCTCGGGTGTGTGTGGTGGCTGCGAAGGC

Sbjct: 3043 CGTAGAGAGGGGGCACTCCGGGAGGACCGTAGCTCGGGTGTGTGTGGTGGCTGCGAAGGC 3102

Query: 1614 TGCTGGCCAGGGAATGGTGGCTGCGGTGGCTGTCTGCTGGCCGCGGGCCCTGAGCGCT 1555  
TGCTGGCCAGGGAATGGTGGCTGCGGTGGCTGTCTGCTGGCCGCGGGCCCTGAGCGCT

Sbjct: 3103 TGCTGGCCAGGGAATGGTGGCTGCGGTGGCTGTCTGCTGGCCGCGGGCCCTGAGCGCT 3162

Query: 1554 CGCTGGGGGCCCCGCTCCCGCGGCCCGCAGGCTGCTGCGTGTGGTGTGGTCCGATTCCG 1495  
CGCTGGGGGCCCCGCTCCCGCGGCCCGCAGGCTGCTGCGTGTGGTGTGGTCCGATTCCG  
Sbjct: 3163 CGCTGGGGGCCCCGCTCCCGCGGCCCGCAGGCTGCTGCGTGTGGTGTGGTCCGATTCCG 3222

Query: 1494 TGCCGCTGCCCCCGGACTTGGAGTCCCCGGCCCTCGGGTCTCTCTCTCTCTCCGATCCG 1435  
TGCCGCTGCCCCCGGACTTGGAGTCCCCGGCCCTCGGGTCTCTCTCTCTCTCCGATCCG  
Sbjct: 3223 TGCCGCTGCCCCCGGACTTGGAGTCCCCGGCCCTCGGGTCTCTCTCTCTCTCCGATCCG 3282

Query: 1434 TGCCGCTACGGTGGAGCCACTGCTCCGACTGCCTTCGCTGTGCTGACTGCTGGCGCTGC 1375  
TGCCGCTACGGTGGAGCCACTGCTCCGACTGCCTTCGCTGTGCTGACTGCTGGCGCTGC  
Sbjct: 3283 TGCCGCTACGGTGGAGCCACTGCTCCGACTGCCTTCGCTGTGCTGACTGCTGGCGCTGC 3342

Query: 1374 CCCC GCCGTAGCTGTAGCCAGCTCCGGGAAGCCCGGGTGC CGGTGTGATGGGTGCGGGG 1315  
CCCCGCCGTAGCTGTAGCCAGCTCCGGGAAGCCCGGGTGC CGGTGTGATGGGTGCGGGG  
Sbjct: 3343 CCCC GCCGTAGCTGTAGCCAGCTCCGGGAAGCCCGGGTGC CGGTGTGATGGGTGCGGGG 3402

Query: 1314 GTGGCGGGTACTGTTACGGGAAGCCATGGGCCAAGGGGCGGCCCGGGTGC GCGCAAG 1255  
GTGGCGGGTACTGTTACGGGAAGCCATGGGCCAAGGGGCGGCCCGGGTGC GCGCAAG  
Sbjct: 3403 GTGGCGGGTACTGTTACGGGAAGCCATGGGCCAAGGGGCGGCCCGGGTGC GCGCAAG 3462

Query: 1254 GGGCCAGTGTGTCTCTGGTCAGAGGCGCCACTGGAGCCATCGTATGCTGGAGAGACAGGT 1195  
GGGCCAGTGTGTCTCTGGTCAGAGGCGCCACTGGAGCCATCGTATGCTGGAGAGACAGGT  
Sbjct: 3463 GGGCCAGTGTGTCTCTGGTCAGAGGCGCCACTGGAGCCATCGTATGCTGGAGAGACAGGT 3522

Query: 1194 TGGCCATGTTGGCGCAGAGGTACCGAAGATGTAGTAGCACTGCTCGGAGAAAGGTGATCT 1135  
TGGCCATGTTGGCGCAGAGGTACCGAAGATGTAGTAGCACTGCTCGGAGAAAGGTGATCT  
Sbjct: 3523 TGGCCATGTTGGCGCAGAGGTACCGAAGATGTAGTAGCACTGCTCGGAGAAAGGTGATCT 3582

Query: 1134 TGTGACGGTATGGCGGATGAAGCCAGCTTTTACAGAGTTGCTGGCATACTTGGCGGCC 1075  
TGTGACGGTATGGCGGATGAAGCCAGCTTTTACAGAGTTGCTGGCATACTTGGCGGCC  
Sbjct: 3583 TGTGACGGTATGGCGGATGAAGCCAGCTTTTACAGAGTTGCTGGCATACTTGGCGGCC 3642

Query: 1074 CCTCCGGTCCGTGAAGCCTTCCACATTGTGGTACAGCCAGTCCACCACATCTGAGCCGA 1015  
CCTCCGGTCCGTGAAGCCTTCCACATTGTGGTACAGCCAGTCCACCACATCTGAGCCGA  
Sbjct: 3643 CCTCCGGTCCGTGAAGCCTTCCACATTGTGGTACAGCCAGTCCACCACATCTGAGCCGA 3702

Query: 1014 TGAAAGCATTAGGGATGGTAATCTTGAGCCACATGCGGTACGGACCTCCAACCTGTATT 955  
TGAAAGCATTAGGGATGGTAATCTTGAGCCACATGCGGTACGGACCTCCAACCTGTATT  
Sbjct: 3703 TGAAAGCATTAGGGATGGTAATCTTGAGCCACATGCGGTACGGACCTCCAACCTGTATT 3762

Query: 954 CAGGGGAGGCCATGGCTTTTACGATGGCAGCCATGTCACTGTGGATGGACAAGTGGAAAT 895  
CAGGGGAGGCCATGGCTTTTACGATGGCAGCCATGTCACTGTGGATGGACAAGTGGAAAT  
Sbjct: 3763 CAGGGGAGGCCATGGCTTTTACGATGGCAGCCATGTCACTGTGGATGGACAAGTGGAAAT 3822

Query: 894 CGTCTAGGCGCTCTGTGTACGGGATGGAACGTGGTATGGAGGAGCTGGTGGAGGTGATGG 835  
CGTCTAGGCGCTCTGTGTACGGGATGGAACGTGGTATGGAGGAGCTGGTGGAGGTGATGG  
Sbjct: 3823 CGTCTAGGCGCTCTGTGTACGGGATGGAACGTGGTATGGAGGAGCTGGTGGAGGTGATGG 3882

Query: 834 TGCTCAAGAAAGTTATATTTTCCATGTTGAGAGTGACCGTATGATGTGATGACATGG 775  
TGCTCA G AG T T C C T T A G G G A G T G T G T G G C A G  
Sbjct: 3883 TGCTCAGGAGGGGC -TCATGCCGTATGCAG -G -GAAGGTGCCGTATGGCTG -CAGTG 3938

Query: 774 TGG -AGTCCGTGAT -GCTGC -TGAA -GGACGAGGACCG -CTCAATCC -GAGAAACCTTCT 721  
TGG AG CC G GC G T AA GG C GGA G CTC TCC G G G AA T  
Sbjct: 3939 TGGGAGAGCCAGGCCGAGGGTCAATGGCC -GGATGGGCTCCTCTCGGCAATGTGAA 3997

Query: 720 GCTTCCGCGCCGCCCTTGTGTCTTCTCATAGGCGTAGGCACTGCTCTGTTCTGTGG 661



Sbjct: 3998 GC CC C G G C TTG GTC CA GGC AG CA G TG C GG  
GCAACAC-GTGGAC--TTGGGTCCCAGCACTTGGCTACAGTCAGGTTGATGGGCCCCGG 4054

Query: 660 AGCTGTGAACCTGCTGGTGGAGTCATCCTCATCTGAGTCAA-AGAAGCTGGTGGTCTCC 602  
TG TG AC CT G AGT A CC A CTG TC A A CT TG TCCT

Sbjct: 4055 TT-TG-TGCACAATCTCCGCACT-ACCCGGA-CTGCATCTCATTA-CTCATGTCTCA 4109

Query: 601 A-GCTCACTGCTCATAAGGGTGGATGAGC-TATCA--TAACCCCTG-GTCC-TC-GCCG 549  
A G T A T CTC T TG A A C TATC T C C TG GTCC TC GC G

Sbjct: 4110 AAGTTGA-T-CTCGTTTACCTGTAACAACATATCTCTGGCTCGATGCGTCCATCAGCAG 4167

Query: 548 CGSTTCCCCCTTCGCACTTCCATTAGCCGG-GTTGCATGCTCTGGGCCATCCCTC-CGG 491  
CC CCCC C T AT AGCC GT G ATGC C G CA C C C CG

Sbjct: 4168 CCAAGCCCCACCTTTCATG-ATAGAGCCAATGTAG-ATGC-CGCCGTACCAACGCTCGT 4224

Query: 490 CGTGGCCGCCCTCGCTGGGCAGACACAAAGA 459  
G CCCC C TGG AGA CC AAGA

Sbjct: 4225 TGCTTTGGCCCAATAG--AGATGCCCAAGA 4254

Score = 2162 (324.4 bits), Expect = 1.2e-182, Sum P(3) = 1.2e-182  
Identities = 466/486 (95%), Positives = 466/486 (95%), Strand = Minus / Plus

Query: 840 TGATGGTGCTCAAGAAGTTATATTTTTCATGTTGAGAGTGACCGTGATGATGTTGAGTG 781  
TG G TGC CAAGAAGTTATATTTTTCATGTTGAGAGTGACCGTGATGATGTTGAGTG

Sbjct: 4240 TGGAGATGCCCAAGAAGTTATATTTTTCATGTTGAGAGTGACCGTGATGATGTTGAGTG 4299

Query: 780 ACATGGTGGAGTCCGTGATGCTGCTGAAGGACGAGGACCGCTCAATCCGAGAAACCTTCT 721  
ACATGGTGGAGTCCGTGATGCTGCTGAAGGACGAGGACCGCTCAATCCGAGAAACCTTCT

Sbjct: 4300 ACATGGTGGAGTCCGTGATGCTGCTGAAGGACGAGGACCGCTCAATCCGAGAAACCTTCT 4359

Query: 720 GCTTCCGCCGCCGCCGCTTGTGTCTTCTCATCAGGCGTGAGGCACCTGCTCTGTTCTGTGG 661  
GCTTCCGCCGCCGCCGCTTGTGTCTTCTCATCAGGCGTGAGGCACCTGCTCTGTTCTGTGG

Sbjct: 4360 GCTTCCGCCGCCGCCGCTTGTGTCTTCTCATCAGGCGTGAGGCACCTGCTCTGTTCTGTGG 4419

Query: 660 AGCTGCTGAACCTGCTGGTGGAGTCATCCTCATCTGAGTCAAAGAAGCTGGTGGTCTCCA 601  
AGCTGCTGAACCTGCTGGTGGAGTCATCCTCATCTGAGTCAAAGAAGCTGGTGGTCTCCA

Sbjct: 4420 AGCTGCTGAACCTGCTGGTGGAGTCATCCTCATCTGAGTCAAAGAAGCTGGTGGTCTCCA 4479

Query: 600 GCTCACTGCTCATAAGGGTGGATGAGCTATCATAACCCCTGGTCTCGCCGCCGTTCC 541  
GCTCACTGCTCATAAGGGTGGATGAGCTATCATAACCCCTGGTCTCGCCGCCGTTCC

Sbjct: 4480 GCTCACTGCTCATAAGGGTGGATGAGCTATCATAACCCCTGGTCTCGCCGCCGTTCC 4539

Query: 540 CCTTCGCACTTCCATTAGCCGGTTGCATGCTCTGGGCCATCCCTCCGGCGTGGCCGCC 481  
CCTTCGCACTTCCATTAGCCGGTTGCATGCTCTGGGCCATCCCTCCGGCGTGGCCGCC

Sbjct: 4540 CCTTCGCACTTCCATTAGCCGGTTGCATGCTCTGGGCCATCCCTCCGGCGTGGCCGCC 4599

Query: 480 CTCGCTGGGCAGACACCAAGAGTCCGTCCTGCTGTCATTGTCCAGGTTCTCTGGCTGC 421  
CTCGCTGGGCAGACACCAAGAGTCCGTCCTGCTGTCATTGTCCAGGTTCTCTGGCTGC

Sbjct: 4600 CTCGCTGGGCAGACACCAAGAGTCCGTCCTGCTGTCATTGTCCAGGTTCTCTGGCTGC 4659

Query: 420 CCCCACCAAGCATGAGGG-TG--G--AAG-GATGGG-GGTCCGGAGTCCCCGATGCCTCCC 368  
CCCCACCAAGCATGAGGG TG G AAG GATGG GG GGG GTC GA G C CCC

Sbjct: 4660 CCCCACCAAGCATGAGGGTGCAGGGAAGAGATGGAAGGATGGGGGTCCG-GA-GTC-CCC 4716

Query: 367 G-TGGCTCC 359  
G TGC CTC

Sbjct: 4717 GATGC-CTCC 4725

Score = 2018 (302.8 bits), Expect = 0.0, Sum P(2) = 0.0  
Identities = 408/412 (99%), Positives = 408/412 (99%), Strand = Minus / Plus

```

Query:   412 GCATGAGGGTGAAGGATGGGGTCGGGAGTCCCGATGCCCTCCGCTCCATAGGT 353
          G A GAG   TGAAGGATGGGGTCGGGAGTCCCGATGCCCTCCGCTCCATAGGT
Sbjct:  4683 GGAAGAGA-TGAAGGATGGGGTCGGGAGTCCCGATGCCCTCCGCTCCATAGGT 4741

Query:   352 GGTGGCAGCTCCGATGGGTATACGACACAGAAGGGGGCTGGGCTGGGTGTGAGCCCTCA 293
          GGTGGCAGCTCCGATGGGTATACGACACAGAAGGGGGCTGGGCTGGGTGTGAGCCCTCA
Sbjct:  4742 GGTGGCAGCTCCGATGGGTATACGACACAGAAGGGGGCTGGGCTGGGTGTGAGCCCTCA 4801

Query:   292 GCTGACACCAGCCAGGACACACCACCGGCCATGAAGCATGGTAGCTTGGCATTGTCAATCC 233
          GCTGACACCAGCCAGGACACACCACCGGCCATGAAGCATGGTAGCTTGGCATTGTCAATCC
Sbjct:  4802 GCTGACACCAGCCAGGACACACCACCGGCCATGAAGCATGGTAGCTTGGCATTGTCAATCC 4861

Query:   232 GAGATCTCCTCCTTACCACCTCCGAAATCGTCCTCATAGACTTGAAGAAGAAGCTTATAG 173
          GAGATCTCCTCCTTACCACCTCCGAAATCGTCCTCATAGACTTGAAGAAGAAGCTTATAG
Sbjct:  4862 GAGATCTCCTCCTTACCACCTCCGAAATCGTCCTCATAGACTTGAAGAAGAAGCTTATAG 4921

Query:   172 CTGGGTCGCTGCAAAACGCCCTTAAAGTCCGCCAAGGTGACGCGCTCGGCGGGCAGGGGC 113
          CTGGGTCGCTGCAAAACGCCCTTAAAGTCCGCCAAGGTGACGCGCTCGGCGGGCAGGGGC
Sbjct:  4922 CTGGGTCGCTGCAAAACGCCCTTAAAGTCCGCCAAGGTGACGCGCTCGGCGGGCAGGGGC 4981

Query:   112 AGCTTCACAAGGTACGGCGTCTCCTCGCCCATCCAAGTGGTAGATGATCTTGGCTCGCCCC 53
          AGCTTCACAAGGTACGGCGTCTCCTCGCCCATCCAAGTGGTAGATGATCTTGGCTCGCCCC
Sbjct:  4982 AGCTTCACAAGGTACGGCGTCTCCTCGCCCATCCAAGTGGTAGATGATCTTGGCTCGCCCC 5041

Query:   52  ATGGCTCTGGCCTCGGGCCCGGCGGCCCGCGCGCGGCGCTGCTCGGCGGCCG 1
          ATGGCTCTGGCCTCGGGCCCGGCGGCCCGCGCGCGGCGCTGCTCGGCGGCCG
Sbjct:  5042 ATGGCTCTGGCCTCGGGCCCGGCGGCCCGCGCGCGGCGCTGCTCGGCGGCCG 5093

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Score = 145 (21.8 bits), Expect = 1.2e-182, Sum P(3) = 1.2e-182  
Identities = 49/68 (72%), Positives = 49/68 (72%), Strand = Minus / Plus

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Query:  1561 GAGCGCTCGCTGGGCGCCCGCTCCCGCGGCCCGCGCAGGCTGCTGCGTGTGGTGTGTC 1502
          GAG GCT   GGC CCC CTCCC C CC CGCAGGCTGC C GTGGT G TCC
Sbjct:  1797 GAGAGCTAAAGAGGCCCCC-CTCCC-C--CCCGCGCAGGCTGCCACACGTGGTGGCATCC 1852

Query:  1501 GATTCTGCT 1494
          GATTCT CT
Sbjct:  1853 GATTCTCT 1860

```

>s3aq:220118507 , 2070 bp.  
Length = 2070

Plus Strand HSPs:

Score = 5677 (851.8 bits), Expect = 2.4e-251, P = 2.4e-251  
Identities = 1329/1501 (88%), Positives = 1329/1501 (88%), Strand = Plus / Plus

```

Query:  459 TCTTTGGTGTCTGCCAGGAGGGCGGCCACGCG-GAGGGATGGCCAGAGCATGCAAC 517
          TCTT GG TCT CC A G GGGC C CGCGAGCTGC C GGCAT C AC
Sbjct:  558 TCTTGGGCATCT-CC-ATTGTGGGCCAAAGCAACGACGTGGTGACGGGC-GCAT-CTAC 613

Query:  518 CC-GGCTAAATGGAACTGCGAAGGGGAACGCGGCG-GA-GGAC-CAGGGGTATGA-- 571

```

GGCT AT G G GGG GGC GC GA GGAC CA G G A GA  
Sbjct: 614 ATTGGCTCTATCATGAAGGGT-GGGGCCGTGGCTGCTGATGGACGCATCGAGCCAGAGA 672

Query: 572 TA-GCTCATCCACCCCTTATGAGCAGTGAGCTG-GAGACCACCAGCTCTTCTT-GACTCAGA 628  
TA G T T CA A GAG A T A CT GAGA CA AG T T G CA GAG  
Sbjct: 673 TATGTTGTTACAGGTAAACGAG-A-TCAACTTTGAGAACATGAG-TAATGACGATGCAG- 728

Query: 629 TGAGGATGACTCCACCAGCAGGTTTACGAGCTCCACAGAACAGAGCAGTGCCTCACGCCCT 688  
T GG T ACT C AG GT CA CA CC G CA C TG CT GCC  
Sbjct: 729 TCCGGT-ACTGCGGGAGATTGTGCA-CAAA-CCGGGGCCCATCACCTGACTGTAGGCA 785

Query: 689 GATGAGAAGACACAAGCGCGCGCGGAAGCAGAAGTTTCTC-GGATTGAGCGG-TCC 746  
TG GAC CAAG C CG GG GC A TT C C GGA GAGC TCC  
Sbjct: 786 AGTGCTGGGACCCAAGTC-CA-CGT-GGTTGCTTACATTGCCACGAGGACGAGCCCATC 842

Query: 747 TCGTCC-TTCAGCA-GCATCACGA-CTCCACCA-TGTCACCTCAACATCATCAGGTCAC 802  
G CC TT A C GC C GG CTCC CA TG CA A C CAC TC C  
Sbjct: 843 G-GCCCATTGACCCCTGGCGCCTGGGTCTCCCACTG-CAGCCATGACCGGCACCTTCCC 900

Query: 803 TCTCAACATGGAAAAATATAACTTCTTGAGCACCATCACCTCCACCAGCTCCTCCATCAC 862  
T CA A GG A A CT C TGAGCACCATCACCTCCACCAGCTCCTCCATCAC  
Sbjct: 901 TG-CAT-ACGSCATGAGCCC-CTCCCTGAGCACCATCACCTCCACCAGCTCCTCCATCAC 957

Query: 863 CAGTTCCATCCCTGACACAGAGCGCCTAGACGACTTCCACTTGTCCATCCAAGTGACAT 922  
CAGTTCCATCCCTGACACAGAGCGCCTAGACGACTTCCACTTGTCCATCCAAGTGACAT  
Sbjct: 958 CAGTTCCATCCCTGACACAGAGCGCCTAGACGACTTCCACTTGTCCATCCAAGTGACAT 1017

Query: 923 GGCTGCCATCGTAAAGCCATGGCCTCCCTGAAATCAGGGTGGAGGTCCGTGACCGCAT 982  
GGCTGCCATCGTAAAGCCATGGCCTCCCTGAAATCAGGGTGGAGGTCCGTGACCGCAT  
Sbjct: 1018 GGCTGCCATCGTAAAGCCATGGCCTCCCTGAAATCAGGGTGGAGGTCCGTGACCGCAT 1077

Query: 983 GTGGCTCAAGATTACCATCCCTAATGCTTTCATCGGCTCAGATGTGGTGGACTGGCTGTA 1042  
GTGGCTCAAGATTACCATCCCTAATGCTTTCATCGGCTCAGATGTGGTGGACTGGCTGTA  
Sbjct: 1078 GTGGCTCAAGATTACCATCCCTAATGCTTTCATCGGCTCAGATGTGGTGGACTGGCTGTA 1137

Query: 1043 CCACAATGTGGAAGGCTTCACGACCGGAGGGAGGCCAGTATGTCAGCAACCTGCT 1102  
CCACAATGTGGAAGGCTTCACGACCGGAGGGAGGCCAGTATGTCAGCAACCTGCT  
Sbjct: 1138 CCACAATGTGGAAGGCTTCACGACCGGAGGGAGGCCAGTATGTCAGCAACCTGCT 1197

Query: 1103 GAAAGCTGGCTTTCATCCGCCATACCGTCAACAAGATCACCTTCTCCGAGCAGTGCTACTA 1162  
GAAAGCTGGCTTTCATCCGCCATACCGTCAACAAGATCACCTTCTCCGAGCAGTGCTACTA  
Sbjct: 1198 GAAAGCTGGCTTTCATCCGCCATACCGTCAACAAGATCACCTTCTCCGAGCAGTGCTACTA 1257

Query: 1163 CATCTTCGGTGACCTCTGCGGCAACATGGCCAACTGTCTCTCCACGATACAGATGGCTC 1222  
CATCTTCGGTGACCTCTGCGGCAACATGGCCAACTGTCTCTCCACGATACAGATGGCTC  
Sbjct: 1258 CATCTTCGGTGACCTCTGCGGCAACATGGCCAACTGTCTCTCCACGATACAGATGGCTC 1317

Query: 1223 CAGTGGCGCCTCTGACCAGGACACACTGGCCCCCTTTCGCGCACCCGGGGCGCGCCCTTG 1282  
CAGTGGCGCCTCTGACCAGGACACACTGGCCCCCTTTCGCGCACCCGGGGCGCGCCCTTG  
Sbjct: 1318 CAGTGGCGCCTCTGACCAGGACACACTGGCCCCCTTTCGCGCACCCGGGGCGCGCCCTTG 1377

Query: 1283 GCCCATGGCTTTCCCGTACCAAGTACCCGCCACCCCGCACCCATACAACCCGCACCCGGG 1342  
GCCCATGGCTTTCCCGTACCAAGTACCCGCCACCCCGCACCCATACAACCCGCACCCGGG  
Sbjct: 1378 GCCCATGGCTTTCCCGTACCAAGTACCCGCCACCCCGCACCCATACAACCCGCACCCGGG 1437

Query: 1343 CTTCCCGGAGCTGGGCTACAGCTACGGCGGGGGCAGCGCCAGCAGTCAGCACAGCGAAGG 1402  
CTTCCCGGAGCTGGGCTACAGCTACGGCGGGGGCAGCGCCAGCAGTCAGCACAGCGAAGG

Sbjct: 1438 CTTCCCGAGCTGGGCTACAGCTACGGCGGGGCGAGCGCAGCAGTCAGCACAGCGAAG 1497

Query: 1403 CAGTCGGAGCAGTGGCTCCAACCGTAGCGGCAGCGATCGGAGGAAGGAGAAGACCCGAA 1462  
CAGTCGGAGCAGTGGCTCCAACCGTAGCGGCAGCGATCGGAGGAAGGAGAAGACCCGAA

Sbjct: 1498 CAGTCGGAGCAGTGGCTCCAACCGTAGCGGCAGCGATCGGAGGAAGGAGAAGACCCGAA 1557

Query: 1463 GGCOCGGGACTTCCAAGTCCGGGGGCGAGCGGACGGAATCGGACCACACACGCGCAGCAG 1522  
GGCOCGGGACTTCCAAGTCCGGGGGCGAGCGGACGGAATCGGACCACACACGCGCAGCAG

Sbjct: 1558 GGCOCGGGACTTCCAAGTCCGGGGGCGAGCGGACGGAATCGGACCACACACGCGCAGCAG 1617

Query: 1523 CCTCGGGGGCCGCGGGAGCGGGGCCGCCAGCGAGCGCTCAGGGCCGGCGGCCAGCGAGCA 1582  
CCTCGGGGGCCGCGGGAGCGGGGCCGCCAGCGAGCGCTCAGGGCCGGCGGCCAGCGAGCA

Sbjct: 1618 CCTCGGGGGCCGCGGGAGCGGGGCCGCCAGCGAGCGCTCAGGGCCGGCGGCCAGCGAGCA 1677

Query: 1583 CAGCCACCCGAGCCACCATTCCCTGGCCAGCAGCCTTCGCGAGCCACCACACACCCCGAG 1642  
CAGCCACCCGAGCCACCATTCCCTGGCCAGCAGCCTTCGCGAGCCACCACACACCCCGAG

Sbjct: 1678 CAGCCACCCGAGCCACCATTCCCTGGCCAGCAGCCTTCGCGAGCCACCACACACCCCGAG 1737

Query: 1643 CTACGGTCTCTCCCGGAGTGCCCCCTCTCTACGGCCCCCCCATTGCTGATGATGCCCCCGCC 1702  
CTACGGTCTCTCCCGGAGTGCCCCCTCTCTACGGCCCCCCCATTGCTGATGATGCCCCCGCC

Sbjct: 1738 CTACGGTCTCTCCCGGAGTGCCCCCTCTCTACGGCCCCCCCATTGCTGATGATGCCCCCGCC 1797

Query: 1703 GCCCGCGGCCATGGGGCCCCAGGAGCCCCCTCGGGCGCGGACCTGGGCTCAGTGCCCCC 1762  
GCCCGCGGCCATGGGGCCCCAGGAGCCCCCTCGGGCGCGGACCTGGGCTCAGTGCCCCC

Sbjct: 1798 GCCCGCGGCCATGGGGCCCCAGGAGCCCCCTCGGGCGCGGACCTGGGCTCAGTGCCCCC 1857

Query: 1763 GGAACCTGACCGCCAGCAGACAGTCTTCGCGATGGCCATGGGAAACCCCAAGTGAGTTCCT 1822  
GGAACCTGACCGCCAGCAGACAGTCTTCGCGATGGCCATGGGAAACCCCAAGTGAGTTCCT

Sbjct: 1858 GGAACCTGACCGCCAGCAGACAGTCTTCGCGATGGCCATGGGAAACCCCAAGTGAGTTCCT 1917

Query: 1823 TGTGGATGTGATGTGAGCAGGGCCCCCTCCCCAGCTCCATTCCGCTCCCAACCCAGCCGG 1882  
TGTGGATGTGATGTGAGCAGGGCCCCCTCCCCAGCTCCATTCCGCTCCCAACCCAGCCGG

Sbjct: 1918 TGTGGATGTGATGTGAGCAGGGCCCCCTCCCCAGCTCCATTCCGCTCCCAACCCAGCCGG 1977

Query: 1883 CTGCGTCTCTCTCCATCCGTCGGTCTTTTACTTTGCTGGTACCTGAAAGGGAAT 1942  
CTGCGTCTCTCTCCATCCGTCGGTCTTTTACTTTGCTGGTACCTGAAAGGGAAT

Sbjct: 1978 CTGCGTCTCTCTCCATCCGTCGGTCTTTTACTTTGCTGGTACCTGAAAGGGAAT 2037

Query: 1943 AAAAGGAACATAATCCA 1959  
AAAAGGAACATAATCCA

Sbjct: 2038 AAAAGGAACATAATCCA 2054

Score = 992 (148.8 bits), Expect = 7.2e-73, Sum P(2) = 7.2e-73

Identities = 200/202 (99%), Positives = 200/202 (99%), Strand = Plus / Plus

Query: 450 GAGACGGACTCTTTGGTGTCTGCCAGCGAGGGGGGCCACGCGGAGGGATGGCCACAG 509  
|||||

Sbjct: 1 GAGACGGACTCTTTGGTGTCTGCCAGCGAGGCGGCCACGCGGAGGGATGGCCACAG 60

Query: 510 CATGCAACCCGGCTAAATGGAACATGCGAAGGGGGAACGCGCGGAGGACAGGGGGTTAT 569  
|||||

Sbjct: 61 CATGCAACCCGGCTAAATGGAACATGCGAAGGGGGAACGCGCGGAGGAAACAGGGGGTTAT 120

Query: 570 GATAGCTCATCCACCCCTTATGAGCAGTGAGCTGGAGACCACAGCTTCTTTGACTCAGAT 629  
|||||

Sbjct: 121 GATAGCTCATCCACCCCTTATGAGCAGTGAGCTGGAGACCACAGCTTCTTTGACTCAGAT 180

Query: 630 GAGGATGACTCCACCAGCAGGT 651

|||||

Sbjct: 181 GAGGATGACTCCACCAGCAGGT 202

Score = 940 (141.0 bits), Expect = 7.2e-73, Sum P(2) = 7.2e-73

Identities = 194/200 (97%), Positives = 194/200 (97%), Strand = Plus / Plus

Query: 641 CACCAGCAGGTTTCAGCAGCTCCACAGAACAGAGCAGTGCCTCAGCGCTGATGAGAAGACA 700

Sbjct: 374 C CCA CAGGTTTCAGCAGCTCCACAGAACAGAGCAGTGCCTCAGCGCTGATGAGAAGACA 432

Query: 701 CAAGCGGCGGCGGGAAGCAGAAGGTTTCTCGGATTGAGCGGTCTCGTCCCTTCAGCAG 760

Sbjct: 433 CAAGCGGCGGCGGGAAGCAGAAGGTTTCTCGGATTGAGCGGTCTCGTCCCTTCAGCAG 492

Query: 761 CATCAGGACTCCACCATTGTCACTCAACATCATCAGGTCACCTCTCAACATGGAAAAATA 820

Sbjct: 493 CATCAGGACTCCACCATTGTCACTCAACATCATCAGGTCACCTCTCAACATGGAAAAATA 552

Query: 821 TAACTTCTTGAGCACCATCA 840

Sbjct: 553 TAACTTCTTG GCA C CA

Sbjct: 553 TAACTTCTTGGGCATCTCCA 572

&gt;s3aq:220119318 , 873 bp.

Length = 873

Plus Strand HSPs:

Score = 4279 (642.0 bits), Expect = 7.8e-188, P = 7.8e-188

Identities = 865/872 (99%), Positives = 865/872 (99%), Strand = Plus / Plus

Query: 362 GCGCAGGGAGGCATCGGGGACTCCCGACCCCATCCTTCCACCCCTCATGCTGGTGGGG 421

Sbjct: 4 GCGCAGGGAGGCATCGGGGACTCCCGACCCCATCCTTCCACCCCTCATGCTGGTGGGG 62

Query: 422 CAGCCAGGAGAACTGGACAATGACACAGAGACGGACTCTTTGGTGTCTGCCACGCGAGG 481

Sbjct: 63 CAGCCAGGAGAACTGGACAATGACACAGAGACGGACTCTTTGGTGTCTGCCACGCGAGG 122

Query: 482 GCGGCCACGCGGAGGGATGGCCAGAGCATGCAACCCGGCTAAATGGAAGTGCAGAGGG 541

Sbjct: 123 GCGGCCACGCGGAGGGATGGCCAGAGCATGCAACCCGGCTAAATGGAAGTGCAGAGGG 182

Query: 542 GGAACGGCGGCGAGGACACAGGGGGTTATGATAGCTCATCCACCTTATGAGCAGTAGCT 601

Sbjct: 183 GGAACGGCGGCGAGGACACAGGGGGTTATGATAGCTCATCCACCTTATGAGCAGTAGCT 242

Query: 602 GGAGACCACCAGCTTCTTTGACTCAGATGAGGATGACTCCACAGCAGGTTTCAGCAGCTC 661

Sbjct: 243 GGAGACCACCAGCTTCTTTGACTCAGATGAGGATGACTCCACAGCAGGTTTCAGCAGCTC 302

Query: 662 CACAGAACAGAGCAGTGCCTCAGCGCTGATGAGAAGACACAAGCGGCGGGCGGGAAGCA 721

Sbjct: 303 CACAGAACAGAGCAGTGCCTCAGCGCTGATGAGAAGACACAAGCGGCGGGCGGGAAGCA 362

Query: 722 GAAGGTTTCTCGGATTGAGCGGTCTCGTCCCTTCAGCAGCATCAGGACTCCACCATTGTC 781

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      GAAGGTTTCTCGGATTGAGCGGTCCTCGTCTTCAGCAGCATCACGGACTCCACCATTGTC
Sbjct: 363 GAAGGTTTCTCGGATTGAGCGGTCCTCGTCTTCAGCAGCATCACGGACTCCACCATTGTC 422

Query: 782 ACTCAACATCATCACGGTCACTCTCAACATGGAAAAATATAACTTCTTGAGCACCATTAC 841
      ACTCAACATCATCACGGTCACTCTCAACATGGAAAAATATAACTTCTTGAGCACCATTAC
Sbjct: 423 ACTCAACATCATCACGGTCACTCTCAACATGGAAAAATATAACTTCTTGAGCACCATTAC 482

Query: 842 CTCACCAGCTCTCCATCACCAGTTCATCCCTGACACAGAGCGCCTAGACGACTTCCA 901
      CTCACCAGCTCTCCATCACCAGTTCATCCCTGACACAGAGCGCCTAGACGACTTCCA
Sbjct: 483 CTCACCAGCTCTCCATCACCAGTTCATCCCTGACACAGAGCGCCTAGACGACTTCCA 542

Query: 902 CTGTGCCATCCACAGTGACATGGCTGCCATCGTAAAAGCCATGGCTCCCTGAATCAGG 961
      CTGTGCCATCCACAGTGACATGGCTGCCATCGTAAAAGCCATGGCTCCCTGAATCAGG
Sbjct: 543 CTGTGCCATCCACAGTGACATGGCTGCCATCGTAAAAGCCATGGCTCCCTGAATCAGG 602

Query: 962 GTTGAGGTTCCGTGACCGCATGTGGCTCAAGATTACCATCCCTTAATGCTTTCATCGGCTC 1021
      GTTGAGGTTCCGTGACCGCATGTGGCTCAAGATTACCATCCCTTAATGCTTTCATCGGCTC
Sbjct: 603 GTTGAGGTTCCGTGACCGCATGTGGCTCAAGATTACCATCCCTTAATGCTTTCATCGGCTC 662

Query: 1022 AGATGTGGTGGACTGGCTGTACCACAATGTGGAAGGCTTCACGGACCGGAGGGAGGCCCG 1081
      AGATGTGGTGGACTGGCTGTACCACAATGTGGAAGGCTTCACGGACCGGAGGGAGGCCCG
Sbjct: 663 AGATGTGGTGGACTGGCTGTACCACAATGTGGAAGGCTTCACGGACCGGAGGGAGGCCCG 722

Query: 1082 CAAGTATGCCAGCAACCTGCTGAAAGCTGGCTTCATCCGCCATACCGTCAACAAGATCAC 1141
      CAAGTATGCCAGCAACCTGCTGAAAGCTGGCTTCATCCGCCATACCGTCAACAAGATCAC
Sbjct: 723 CAAGTATGCCAGCAACCTGCTGAAAGCTGGCTTCATCCGCCATACCGTCAACAAGATCAC 782

Query: 1142 CTTCTCGAGCAGTGCTACTACATCTTCGGTGACCTCTGCGGCAACATGGCCAACTGTC 1201
      CTTCTCGAGCAGTGCTACTACATCTTCGGTGACCTCTGCGGCAACATGGCCAACTGTC
Sbjct: 783 CTTCTCGAGCAGTGCTACTACATCTTCGGTGACCTCTGCGGCAACATGGCCAACTGTC 842

Query: 1202 TCTCCACGATCAGATGGCTCCAGTGGCGCCT 1233
      TCTCCACGATCAGATG CTCC GTGG GCCT
Sbjct: 843 TCTCCACGATCAGATGCCCTCC-GTGG-GCCT 872

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>s3aq:220118872 , 474 bp.  
Length = 474

Minus Strand HSPs:

Score = 2340 (351.1 bits), Expect = 5.4e-100, P = 5.4e-100  
Identities = 470/473 (99%), Positives = 470/473 (99%), Strand = Minus / Plus

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Query: 1959 TGGATTATTAGTTCCTTTTATTTCCTTTTCAGGTACGACACAAAGTAAAAAGACGGACGGA 1900
      |||
Sbjct: 1 TGGATTATTAGTTCCTTTTATTTCCTTTTCAGGTACGACACAAAGTAAAAAGACGGACGGA 60

Query: 1899 TGGAGAGAGGAACGACGCGGCTGGGGTGGGAGCGGAATGGAGCTGGGGGAGGGGCCCTG 1840
      |||
Sbjct: 61 TGGAGAGAGGAACGACGCGGCTGGGGTGGGAGCGGAATGGAGCTGGGGGAGGGGCCCTG 120

Query: 1839 CTCACATCACAATCCACAAAGAACTCACTGGGGTTTCCCATGGCCATGCGGAAGGACTGTC 1780
      |||
Sbjct: 121 CTCACATCACAATCCACAAAGAACTCACTGGGGTTTCCCATGGCCATGCGGAAGGACTGTC 180

Query: 1779 TGTGCGCGGTCACTCCGGGGGCACTGAGGCCAGGTGCGCGCCCGGAGGGGCTCTGCGG 1720

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|||||
Sbjct: 181 TGCTGGCGGTCAGTTCGCGGGGCACTGAGGCCAGGTCGCGGCCGCGAGGGGCTCCTCGGG 240
Query: 1719 GCCCATGGCCGCGGGCGCGGGGGCATCATCAGCATGGGGGGCCGTAGAGAGGGGGCA 1660
|||||
Sbjct: 241 GCCCATGGCCGCGGGCGCGGGGGCATCATCAGCATGGGGGGCCGTAGANAGGGGGCA 300
Query: 1659 CTCGGGAGGACCGTAGCTCGGGTGTGTGTGGTGGCTGCGAAGGCTGCTGGCCAGGGAAT 1600
|||||
Sbjct: 301 CTCGGGAGGACCGTAGCTCGGGTGTGTGTGGTGGCTGCGAAGGCTGCTGGCCAGGGAAT 360
Query: 1599 GGTGGCTGCGGTGGCTGTGCTCGCTGGCCGCGGCCCTGAGCGCTCGCTGGCGCCCGCT 1540
|||||
Sbjct: 361 GGTGGCTGCGGTGGCTGTGCTCGCTGGCGCGGCCCTGAGCGCTCGCTGGCGCCCGCT 420
Query: 1539 CCCGCGGCCCCCGCAGGCTGCTGCGTGTGGTGTGGTCCGATTGCTGCGCGTG 1487
|||||
Sbjct: 421 CCCGCGGACCCCGCAGGCTGCTGCGTGTGGTGTGGTCCGATTGCTGCTGCTG 473
```

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>s3aq:220119337 , 373 bp.
Length = 373
```

Minus Strand HSPs:

Score = 1849 (277.4 bits), Expect = 1.0e-77, P = 1.0e-77  
Identities = 371/373 (99%), Positives = 371/373 (99%), Strand = Minus / Plus

```
Query: 1948 CCTTTTATTTCCTTTTCAGGTACCGACAAAGTAAAAAGACGGACGGATGGAGAGAGGA 1889
|||||
Sbjct: 1 CCTTTTATTTCCTTTTCAGGTACCGACAAAGTAAAAAGACGGACGGATGGAGAGAGGA 60
Query: 1888 ACGCAGCCGGCTGGGGTGGGAGCGGAATGGAGCTGGGGGAGGGGCCCTGCTCACATCACA 1829
|||||
Sbjct: 61 ACGCAGCCGGCTGGGGTGGGAGCGGAATGGAGCTGGGGGAGGGGCCCTGCTCACATCACA 120
Query: 1828 TCCACAAGAAGTCACTGGGGTTTCCCATGGCCATGCGGAAGGACTGTCTGCTGGCGGTC 1769
|||||
Sbjct: 121 TCCACAAGAAGTCACTGGGGTTTCCCATGGCCATGCGGAAGGACTGTCTGCTGGCGGTC 180
Query: 1768 AGTTCCGGGGGCACTGAGGCCAGGTCGCGGCCCGGAGGGGCTCCTGGGGGCCCATGGCC 1709
|||||
Sbjct: 181 AGTTCCGGGGGCACTGAGGCCAGGTCGCGGCCCGGAGGGGCTCCTGGGGGCCCATGGCC 240
Query: 1708 GCGGGCGCGGGGGGCATCATCAGCATGGGGGGCCGTAGAGAGGGGGCACTCCGGGGAGGA 1649
|||||
Sbjct: 241 GCGGGCGCGGGGGGCATCATCAGCATGGGGGGCCGTAGAGAGGGGGCACTCCGGGGAGGA 300
Query: 1648 CGTAGCTCGGGTGTGTGTGGTGGCTGCGAAGGCTGCTGGCCAGGGAATGGTGGCTGCGG 1589
|||||
Sbjct: 301 CGTAGCTCGGGTGTGTGTGGTGGCTGCGAAGGCTGCTGGCCAGGGAATGGTGGCTGCGG 360
Query: 1588 TGGCTGTGCTCGC 1576
|||||
Sbjct: 361 TGGCTGTGCTCGC 373
```

>s3aq:220119235 , 625 bp.  
Length = 625

## Minus Strand HSPs:

Score = 1695 (254.3 bits), Expect = 5.6e-71, P = 5.6e-71  
Identities = 415/466 (89%), Positives = 415/466 (89%), Strand = Minus / Plus

Query: 1611 TGGCCA-G-GGAATGG-TGGCTGC-GGTGG-CTGTGCTCGTGGCCGCCG-GCCCTGAGC 1558  
TGGCCA G GGAA G TG CTGC GG GG C GT C GG GGC C G G CC G C  
Sbjct: 160 TGGCCATGCGGAAGGACTGTCTGCTGGCGGTTCAGTTC-CGGGGGCA-CTGAGGCCAGGTC 217

Query: 1557 GCTGCTGGGGCGCCCGCTCCCAGGGCCCGCAGGCTGCTGCGTGTGGTGTGGTCCGATT 1498  
GC C GG G GCTCC G GG CCCC GGC GCTGCGTGTGGTGTGGTCCGATT  
Sbjct: 218 GCGGGCCGGTAGGG-GCTCCTGGGGGCCCAT-GGC-GCTGCGTGTGGTGTGGTCCGATT 274

Query: 1497 CGCTGCCGCTGCCCCCGGACTTGGAGTCCCGGCCCTTCGGGTCTTCTCCTTCCTCCGAT 1438  
CGCTGCCGCTGCCCCCGGACTTGGAGTCCCGGCCCTTCGGGTCTTCTCCTTCCTCCGAT  
Sbjct: 275 CGCTGCCGCTGCCCCCGGACTTGGAGTCCCGGCCCTTCGGGTCTTCTCCTTCCTCCGAT 334

Query: 1437 CGCTGCCGCTACGGTTGGAGCCACTGCTCCGACTGCCTTCG-CTGTGCTGACTGTGG-C 1380  
CGCTGCCGCTACGGTTGGAGCCACTGCTCCGACTGCCTTCG CTGTGCTGACTGTGG C  
Sbjct: 335 CGCTGCCGCTACGGTTGGAGCCACTGCTCCGACTGCCTTCGCTGACTGTGGCG 394

Query: 1379 GCTGCCCCCGCGTAGCTGTAGCCAGCTCCGGGAAGCCGGGTGCGGGTTGTATGGGTG 1320  
GCTGCCCC GCGTAGCTGTAGCCAGCTCCGGGAAGCCGGGTGCGGGTTGTATGGGTG  
Sbjct: 395 GCTGCCCC-GCGTAGCTGTAGCCAGCTCCGGGAAGCCGGGTGCGGGTTGTATGGGTG 452

Query: 1319 CGGGGGTGGGGGTACTGGTACGGGAAGCCATGGGCCAAGGGGCGGCCCGGGTGGCG 1260  
CGGGG TGGCGGGTACTGGTACGGGAAGCCATGGGCCAAGGGGCGGCCCGGGTGGCG  
Sbjct: 453 CGGGG-TGGCGGGTACTGGTACGGGAAGCCATGGGCCAAGGGGCGGCCCGGGTGGCG 511

Query: 1259 CAAAGGGGCCAGTGTGTCTGGTCAGAGGCGCCACTGGAGCCATCGTGATCGTGAGAGA 1200  
CAAAGGGGCCAGTGTGTCTGGTCAGAGGCGCCACTGGAGCCATCGTGATCGTGAGAGA  
Sbjct: 512 CAAAGGGGCCAGTGTGTCTGGTCAGAGGCGCCACTGGAGCCATCGTGATCGTGAGAGA 571

Query: 1199 CAGGTTGGCCATGT-TGCCGCGAGAGTCA-CCGAAGATGTAG-TAGCACTGCTCGGA 1146  
C GGTGGCCATGT TG G AG GG CCG GATG AG TAG CT CGGA  
Sbjct: 572 CGGTTGGCCATGTCTGT-GAAGGGGAGGGCG--GATGGAGTGGTCTTCCCGA 625

Score = 1321 (198.2 bits), Expect = 2.3e-53, P = 2.3e-53  
Identities = 373/460 (81%), Positives = 373/460 (81%), Strand = Minus / Plus

Query: 1958 GGATTTTAGTTCCTTTTATTTCCCTTTTCAGGTACACAGCAAAAGTAAAAA-GACGGACGGA 1900  
GGATTTTAGTTCCTTTTATTTCCCTTTTCAGGTACACAGCAAAAGTAAAAA GACGGACGGA  
Sbjct: 1 GGATTTTAGTTCCTTTTATTTCCCTTTTCAGGTACACAGCAAAAGTAAAAAAGACGGACGGA 60

Query: 1899 TGGAGAGAGGAACGACAGCCGGCTGGGGTGGGAGCGGAATGGAGCTGGGGGAGGGGCCCTG 1840  
TGGAGAGAGGAACGACAGCCGGCTGGGGTGGGAGCGGAATGGAGCTGGGGGAGGGGCCCTG  
Sbjct: 61 TGGAGAGAGGAACGACAGCCGGCTGGGGTGGGAGCGGAATGGAGCTGGGGGAGGGGCCCTG 120

Query: 1839 CTCACATCACATCCACAAAGAACTCACTGGGGTTTCCCATGGCCATGCGGAAGGACTGTC 1780  
CTCACATCACATCCACAAAGAACTCACTGGGGTTTCCCATGGCCATGCGGAAGGACTGTC  
Sbjct: 121 CTCACATCACATCCACAAAGAACTCACTGGGGTTTCCCATGGCCATGCGGAAGGACTGTC 180

Query: 1779 TGCTGSCGCTCAGTTCGGGGGGCACTAGGCCAGGTGCGGGCCCGG-AGGGGCTCTGGG 1721  
TGCTGSCGCTCAGTTCGGGGGGCACTAGGCCAGGTGCGGGCCCGG AGGGGCTCTGGG



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>s3aq:220120226 , 345 bp.
      Length = 345

      Minus Strand HSPs:

      Score = 1313 (197.0 bits), Expect = 7.1e-70, Sum P(2) = 7.1e-70
      Identities = 267/271 (98%), Positives = 267/271 (98%), Strand = Minus / Plus

Query:   496   CTCGCGCGTGGCGGCCCTCGCTGGGCAGACACAAAGAGTCCGCTCTCTGTGTCATTGTGCC 437
          CTCC G G GCGCGC CTCGCTGGGCAGACACAAAGAGTCCGCTCTCTGTGTCATTGTGCC
Sbjct:   76    CTCCTGGG-GGCGCGCTCTCGCTGGGCAGACACAAAGAGTCCGCTCTCTGTGTCATTGTGCC 134

Query:   436   AGGTTCTCCTGGCTGCCCCACCAGCATGAGGGTGGAAAGGATGGGGGTGGGAGTCCCG 377
          AGGTTCTCCTGGCTGCCCCACCAGCATGAGGGTGGAAAGGATGGGGGTGGGAGTCCCGG
Sbjct:   135   AGGTTCTCCTGGCTGCCCCACCAGCATGAGGGTGGAAAGGATGGGGGTGGGAGTCCCGG 194

Query:   376   ATGCCCTCCCGTGCCTCCATAGGTGGTGGCAGCTCCGATGGGTTATCAGCACAGAAGGGG 317
          ATGCCCTCCCGTGCCTCCATAGGTGGTGGCAGCTCCGATGGGTTATCAGCACAGAAGGGG
Sbjct:   195   ATGCCCTCCCGTGCCTCCATAGGTGGTGGCAGCTCCGATGGGTTATCAGCACAGAAGGGG 254

Query:   316   GCTGGGTCTGGGTTGAGAGCCCTCAGCTGACACAGCCAGGACACACCCGGGCCATTGAAG 257
          GCTGGGTCTGGGTTGAGAGCCCTCAGCTGACACAGCCAGGACACACCCGGGCCATTGAAG
Sbjct:   255   GCTGGGTCTGGGTTGAGAGCCCTCAGCTGACACAGCCAGGACACACCCGGGCCATTGAAG 314

Query:   256   CATGGTAGCTTGGCATTTGTATCCGAGATCT 226
          CATGGTAGCTTGGCATTTGTATCCGAGATCT
Sbjct:   315   CATGGTAGCTTGGCATTTGTATCCGAGATCT 345

      Score = 439 (65.9 bits), Expect = 7.1e-70, Sum P(2) = 7.1e-70
      Identities = 97/103 (94%), Positives = 97/103 (94%), Strand = Minus / Plus

Query:   1803  CCATGGCCATGCGGAAGGACTGTCTGCTGGCGGTGAGTCCGGGGGCACTGAGGCCAGGT 1744
          CCATGGCCATGCGGAAGGACTGTCTGCTGGCGGTGAGTCCGGGGGCACTGAGGCCAGGT
Sbjct:   1     CATGCGCATGCGGAAGGACTGTCTGCTGGCGGTGAGTCCGGGGGCACTGAGGCCAGGT 60

Query:   1743  CGCGGGCCGAGGGGCTCCTGGGGGCCCCATGCCGCG-GGGCGG 1701
          CGCGGGCCGAGGGGCTCCTGGGGGCCCC C T C GC GGGC G
          CGCGGGCCGAGGGGCTCCTGGGGGCGCG-TCTC-GCTGGGCG 102
Sbjct:   61    CGCGGGCCGAGGGGCTCCTGGGGGCGCG-TCTC-GCTGGGCG 102

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**Figure 4. ClustalW alignment of CG164330-01 protein with related proteins.**

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CG164330_01  MQETK I I Y H L D G Q E T P Y L V K L P L P A E R V T L A D F K Q V L Q R F S Y K F F F K S M D D F G V V K E E I
DVL3_HUMAN   MQETK I I Y H L D G Q E T P Y L V K L P L P A E R V T L A D F K Q V L Q R F S Y K F F F K S M D D F G V V K E E I
DVL3_MOUSE   MQETK I I Y H L D G Q E T P Y L V K L P L P A E R V T L A D F K Q V L Q R F S Y K F F F K S M D D F G V V K E E I

CG164330_01  S D D N A K L P C F N G R V V S W L V S A E Q S H P D A P F C A D N F S E L P P P M E R T G G I G D S R P P S P H P H
DVL3_HUMAN   S D D N A K L F C F N G R V V S W L V S A E Q S H P D A P F C A D N F S E L P P P M E R T G G I G D S R P P S P H P H
DVL3_MOUSE   S D D N A K L F C F N G R V V S W L V S A E Q S H P D A P F C A D N F S E L P P S M E R T G G I G D S R P P S P H P H

CG164330_01  A G G G S Q Z W L D N D T E T D S L V S A C Q F P R R D C P E H A T R L N G T A L G E R R R P P G G Y S S S T L M
DVL3_HUMAN   A G G G Q E H L D N D T E T D S L V S A C A E P R R D C P E H A T R L N T A K E R R E P P G G Y S S S T L M
DVL3_MOUSE   A S G Q G E N L D N D T E T D S L V S A C R E P R R R D G F E A P L N O T L G E R R R P P G G Y S S S T L M

CG164330_01  S S L E T T S F F D S D E D D S T S R F S S S T E Q S S A S R L M R R H K R R R K Q K V S R I E R S S S F S S I T D
DVL3_HUMAN   S S L E T T S F F D S D E D D S T S R F S S S T E Q S S A S R L M R R H K R R R K Q K V S R I E R S S S F S S I T D
DVL3_MOUSE   S S L E T T S F F D S D E D D S T S R F S S S T E Q S S A S R L M R R H K R R R K Q K V S R I E R S S S F S S I T D

CG164330_01  S T M S L N I I T V T L N M E K Y N F L G I S I V Q G S N E R O D G G I Y I G S I M E G G A Y A A D G R I E P Q D M L L
DVL3_HUMAN   S T M S L N I I T V T L N M E K Y N F L G I S I V Q G S N E R O D G G I Y I G S I M E G G A Y A A D G R I E P Q D M L L
DVL3_MOUSE   S T M S L N I I T V T L N M E K Y N F L G I S I V Q G S N E R O D G G I Y I G S I M E G G A Y A A D G R I E P Q D M L L

CG164330_01  Q V N E I N F E N M S N D D A V R V L R E I V H K P Q P I T L T V A K W D F S P R G C P T L P R S E P I R P I D P A A
DVL3_MOUSE   Q V N E I N F E N M S N D D A V R V L R E I V H K P Q P I T L T V A K W D F S P R G C P T L P R S E P I R P I D P A A

CG164330_01  T I T S T S S I T S S I P D T E R L D D F H L S I H S D M A A I V K A M A
DVL3_HUMAN   W S S H T A A M T G T F P A Y G M S F S L S T I T S T S S I T S S I P D T E R L D D F H L S I H S D M A A I V K A M A
DVL3_MOUSE   W S S H T A A M T G T F P A Y G M S F S L S T I T S T S S I T S S I P D T E R L D D F H L S I H S D M A A I V K A M A

CG164330_01  S P E S G L E V R E R M W L K I T I P N A F I Q S D V V D W L Y H N V G E F T D R R E A R K Y A S N L K A G F I R H T
DVL3_HUMAN   S P E S G L E V R E R M W L K I T I P N A F I Q S D V V D W L Y H N V G E F T D R R E A R K Y A S N L K A G F I R H T
DVL3_MOUSE   S P E S G L E V R E R M W L K I T I P N A F I Q S D V V D W L Y H N V G E F T D R R E A R K Y A S N L K A G F I R H T

CG164330_01  V N K I T F S E Q C Y Y I F O D L C G N M A N L S L H E D G S S G A S D Q D T L A P L P H G A A P W P M A F P Y Q Y
DVL3_MOUSE   V N K I T F S E Q C Y Y I F O D L C G N M A N L S L H E D G S S G A S D Q D T L A P L P H G A A P W P M A F P Y Q Y

CG164330_01  P P P P H P Y N P H G F P P E L Q V S Y G G S A S S Q H S E Q S R S S Q S M R S Q S D R A K E K D P K A G D S K S G G
DVL3_HUMAN   P P P P H P Y N P H G F P P E L Q V S Y G G S A S S Q H S E Q S R S S Q S M R S Q S D R A K E K D P K A G D S K S G G
DVL3_MOUSE   P P P P H P Y N P H G F P P E L Q V S Y G G S A S S Q H S E Q S R S S Q S M R S Q S D R A K E K D P K A G D S K S G G

CG164330_01  S G S E S D H T T R S S L R G F R E R A P S E R S O P A A S E H S H R S H H S L A S S L R S H N T H P S Y P G P V P P
DVL3_HUMAN   S G S E S D H T T R S S L R G F R E R A P S E R S O P A A S E H S H R S H H S L A S S L R S H N T H P S Y P G P V P P
DVL3_MOUSE   S G S E S D H T T R S S L R G F R E R A P S E R S O P A A S E H S H R S H H S L A S S L R S H N T H P S Y P G P V P P

CG164330_01  L Y G P F M L M M P P P F A A M G P F G A P F O R D L A S V P P E L T A S R Q S F R I M A M O N P S E F F V D V M
DVL3_HUMAN   L Y G P F M L M M P P P F A A M G P F G A P F O R D L A S V P P E L T A S R Q S F R I M A M O N P S E F F V D V M
DVL3_MOUSE   L Y G P F M L M M P P P F A A M G P F G A P F O R D L A S V P P E L T A S R Q S F R I M A M O N P S E F F V D V M

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Information for the ClustalW proteins:

Accno	Common Name	Length
CG164330_01	novel Dishevelled-3-like protein	595
DVL3_HUMAN	Segment polarity protein dishevelled homolog DVL-3 (Dishevelled-3) DE (DSH homolog 3).	716
DVL3_MOUSE	Segment polarity protein dishevelled homolog DVL-3 (Dishevelled-3) DE (DSH homolog 3).	716

In the alignment shown above, black outlined amino acid residues indicate residues identically conserved between sequences (i.e., residues that may be required to preserve structural or functional properties); amino acid residues with a gray background are similar to one another

between sequences, possessing comparable physical and/or chemical properties without altering protein structure or function (e.g. the group L, V, I, and M may be considered similar); and amino acid residues with a white background are neither conserved nor similar between sequences.

**Figure 5: PSORT, SignalP and hydropathy results for CuraGen Acc. No. CG164330-01.**

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nucleus --- Certainty=0.7000(Affirmative) < succ>
microbody (peroxisome) --- Certainty=0.4022(Affirmative) < succ>
mitochondrial matrix space --- Certainty=0.1000(Affirmative) < succ>
lysosome (lumen) --- Certainty=0.1000(Affirmative) < succ>

```

Is the sequence a signal peptide?

# Measure	Position	Value	Cutoff	Conclusion
max. C	32	0.087	0.37	NO
max. Y	39	0.053	0.34	NO
max. S	31	0.168	0.88	NO
mean S	1-38	0.070	0.48	NO

